

## References

- Adams RJ, Fuhlbrigge A, Finkelstein JA, Lozano P, Livingston JM, Weiss KB, Weiss ST. Impact of inhaled anti-inflammatory therapy on hospitalization and emergency department visits for children with asthma. *Pediatrics* 2001;107(4):706-11.
- Agertoft L, Pedersen S. Effects of long-term treatment with inhaled budesonide on adult height in children with asthma. *N Engl J Med* 2000;343(15):1064-69.
- Agertoft L, Pedersen S. Effects of long-term treatment with an inhaled corticosteroid on growth and pulmonary function in asthmatic children. *Respir Med* 1994;88(5):373-81.
- Altman LC, Munk Z, Seltzer J, Noonan N, Shingo S, Zhang J, Reiss TF. A placebo-controlled, dose-ranging study of montelukast, a cysteinyl leukotriene-receptor antagonist. Montelukast Asthma Study Group. *J Allergy Clin Immunol* 1998;102(1):50-6.
- Baker JW, Mellon M, Wald J, Welsh M, Cruz-Rivera M, Walon-Bowen K. A multiple-dosing, placebo-controlled study of budesonide inhalation suspension given once or twice daily for treatment of persistent asthma in young children and infants. *Pediatrics* 1999;103:414-21.
- Bisgaard H. Future options for aerosol delivery to children. *Allergy* 1999;54 Suppl 49:97-103.
- Bleecker ER, Welch MJ, Weinstein SF, Kalberg C, Johnson M, Edwards L, Rickard KA. Low-dose inhaled fluticasone propionate versus oral zafirlukast in the treatment of persistent asthma. *J Allergy Clin Immunol* 2000;105(6 Pt 1):1123-9.
- Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.
- Busse W, Raphael GD, Galant S, Kalberg C, Goode-Sellers S, Srebro S, Edwards L, Rickard K. Low-dose fluticasone propionate compared with montelukast for first-line treatment of persistent asthma: A randomized clinical trial. *J Allergy Clin Immunol* 2001;107(3):461-8.
- Castro-Rodriguez JA, Holberg CJ, Wright AL, Martinez FD. A clinical index to define risk of asthma in young children with recurrent wheezing. *Am J Respir Crit Care Med* 2000;162(4 Pt 1):1403-6.
- Childhood Asthma Management Program (CAMP) Research Group. Long-term effects of budesonide or nedocromil in children with asthma. *N Engl J Med* 2000;343(15):1054-63.
- Connett GJ, Warde C, Wooler E, Lenney W. Use of budesonide in severe asthmatics aged 1-3 years. *Arch Dis Child* 1993;69(3):351-5.
- de Blic J, Delacourt C, Le Bourgeois M, Mahut B, Ostinelli J, Caswell C, Scheinmann P. Efficacy of nebulized budesonide in treatment of severe infantile asthma: a double-blind study. *J Allergy Clin Immunol* 1996;98(1):14-20.
- Donahue JG, Weiss ST, Livingston JM, Goetsch MA, Gremmeder DK, Platt R. Inhaled steroids and the risk of hospitalization for asthma. *JAMA* 1997;277(11):887-91.
- DuBuske LM, Grossman J, Dube LM, Swanson LJ, Lancaster JF. Randomized trial of zileuton in patients with moderate asthma: effect of reduced dosing frequency and amounts on pulmonary function and asthma symptoms. Zileuton Study Group. *Am J Manag Care* 1997;3(4):633-40.
- Hoekstra MO, Grol MH, Bouman K, Stijnen T, Koeter GH, Kauffman HF, Gerritsen J. Fluticasone propionate in children with moderate asthma. *Am J Respir Crit Care Med* 1996;154(4 Pt 1):1039-44.
- Hoekstra MO, Grol MH, Hovenga H, Bouman K, Stijnen T, Koeter GH, Gerritsen J, Kauffman HF. Eosinophil and mast cell parameters in children with stable moderate asthma. *Pediatr Allergy Immunol* 1998;9(3):143-9.

Israel E, Cohn J, Dube L, Drazen JM. Effect of treatment with zileuton, a 5-lipoxygenase inhibitor, in patients with asthma. A randomized controlled trial. Zileuton Clinical Trial Group. *JAMA* 1996;275(12):931-6.

Jousson G, Carlsen KH, Blomqvist P. Clinical efficacy of low-dose inhaled budesonide once or twice daily in children with mild asthma not previously treated with steroids. *Eur Respir J* 1998;12(5):1099-104.

Kemp JP, Dockhorn RJ, Shapiro GG, Nguyen HH, Reiss TF, Seidenberg BC, Knorr B. Montelukast once daily inhibits exercise-induced bronchoconstriction in 6- to 14-year-old children with asthma. *J Pediatr* 1998;133(3):424-8.

Kemp JP, Skoner DP, Szeffler SJ, Walton-Brown K, Cruz-Rivera M, Smith JA. Once-daily budesonide inhalation suspension for the treatment of persistent asthma in infants and young children. *Ann Allergy Asthma Immunol* 1999;83(3):231-9.

Knorr B, Franchi LM, Bisgaard H, Vermeulen JH, LeSouef O, Santanello N, Michele TM, Reiss TF, Nguyen HH, Bratton DL. Montelukast, a leukotriene receptor antagonist, for the treatment of persistent asthma in children aged 2 to 5 years. *Pediatrics* 2001;108(3):E48.

Knorr B, Matz J, Bernstein JA, Nguyen H, Seidenberg BC, Reiss TF, Becker A. Montelukast for chronic asthma in 6- to 14-year old children: a randomized double-blind trial. Pediatric Montelukast Study Group. *JAMA* 1998;279(15):1181-6.

Konig P. Evidence for benefits of early intervention with non-steroidal drugs in asthma. *Pediatr Pulmonol Suppl* 1997;15:34-9.

Lazarus SC, Boushey HA, Fahy JV, Chinchilli VM, Lemanske RF Jr, Sorkness CA, Kraft M, Fish JE, Peters SP, Craig T, et al. Asthma Clinical Research Network for the National Heart, Lung, and Blood Institute. Long-acting beta<sub>2</sub>-agonist monotherapy vs. continued therapy with inhaled corticosteroids in patients with persistent asthma: a randomized controlled trial. *JAMA* 2001;285(20):2583-93.

Martinez FD. Viral infections and the development of asthma. *Am J Res Crit Care Med* 1995;151(5):1644-7.

Martinez FD, Wright AL, Taussig LM, Holberg CJ, Halonen M, Morgan WJ. The Group Health Medical Associates. Asthma and wheezing in the first six years of life. *N Engl J Med* 1995;332(3):133-8.

Nathan RA, Bernstein JA, Bielory L, Bonuccelli CM, Calhoun WJ, Galant SP, Hanby LA, Kemp JP, Kylstra JW, Nayak AS, O'Connor JP, Schwartz HJ, Southern DL, Spector SL, Williams PV. Zafirlukast improves asthma symptoms and quality of life in patients with moderate reversible air-flow obstruction. *J Allergy Clin Immunol* 1998;102(6 Pt 1):935-42.

Nielsen KG, Bisgaard H. The effect of inhaled budesonide on symptoms, lung function, and cold air and methacholine responsiveness in 2-5 year old asthmatic children. *Am J Res Crit Care Med* 2000;162:1500-6.

Pearlman DS, Lampl KL, Dowling PJ Jr., Miller CJ, Bonuccelli CM. Effectiveness and tolerability of zafirlukast for the treatment of asthma in children. *Clin Ther* 2000;22(6):732-47.

Petty TL, Rollins DR, Christopher K, Good JT, Oakley R. Cromolyn sodium is effective in adult chronic asthmatics. *Am Rev Respir Dis* 1989;139(3):694-701.

Reed CE, Offord KP, Nelson HS, Li JT, Tinkelman DG. Aerosol beclomethasone dipropionate spray compared with theophylline as primary treatment for chronic mild or moderate persistent asthma. *J Allergy Clin Immunol* 1998;101:14-23.

Schwartz HJ, Petty T, Dube LM, Swanson LJ, Lancaster JE. A randomized controlled trial comparing zileuton with theophylline in moderate asthma. The Zileuton Study Group. *Arch Intern Med* 1998;158(2):141-8.

Storr J, Lenney CA, Lenney W. Nebulized beclomethasone dipropionate in preschool asthma. *Arch Dis Child* 1986;61(3):270-3.

Tasche MJ, Uijen JH, Bernsen RM, de Jongste JC, van der Wouden JC. Inhaled disodium cromoglycate (DSCG) as maintenance therapy in children with asthma: a systematic review. *Thorax* 2000;55(11):913-20.

Tashkin DP, Nathan RA, Howland WC, Minkwitz MC, Simonson SG, Bonuccelli CM. An evaluation of zafirlukast in the treatment of asthma with exploratory subset analyses. *J Allergy Clin Immunol* 1999;103(2 Pt 1):246-54.

Tinkelman DG, Reed CE, Nelson HS, Offord KP. Aerosol beclomethasone dipropionate compared with theophylline as primary treatment of chronic, mild to moderately severe asthma in children. *Pediatrics* 1993;92(1):64-77.

van Essen-Zandvliet EE, Hughes MD, Waalkens HJ, Duiverman EJ, Pocock SJ, Kerrebijn KF. Effects of 22 months of treatment with inhaled corticosteroids and/or beta-2-agonists on lung function, airway responsiveness, and symptoms in children with asthma. The Dutch Chronic Non-specific Lung Disease Study Group. *Am Rev Respir Dis* 1992;146(3):547-54.

Verberne AA, Frost C, Roorda RJ, van der Laag H, Kerrebijn KF. One year treatment with salmeterol compared with beclomethasone in children with asthma. The Dutch Paediatric Asthma Study Group. *Am J Respir Crit Care Med* 1997;156(3 Pt 1):688-95.

Weinberger M. Zafirlukast and cromolyn are effective first-line therapies for child asthma. *Ann Allergy Asthma Immunol* 2000;84(6):638-9.

## Long-Term Management of Asthma in Children: Safety of Inhaled Corticosteroids

### Question

*What are the long-term adverse effects of chronic inhaled corticosteroid use in children on the following outcomes?*

- Vertical growth?
- Bone mineral density (BMD)?
- Ocular toxicity?
- Suppression of adrenal/pituitary axis?

### Summary Answer to the Question

Strong evidence from clinical trials following children for up to 6 years suggests that the use of inhaled corticosteroids at recommended doses does not have long-term, clinically significant, or irreversible effects on any of the outcomes reviewed. Inhaled corticosteroids do improve health outcomes for children with mild or moderate persistent asthma, and the potential but small risk of delayed growth is well balanced by their effectiveness (SRE-Evidence A, B). Updated text is recommended for the EPR-2 incorporating the results of the SRE, but this update does not change the EPR-2 statements.

### Rationale for the Question

Inhaled corticosteroids have been proven to be beneficial in the treatment of mild or moderate persistent asthma in children. Because this class of compounds has the potential for producing adverse side effects, however, a SRE on the potential long-term adverse effects would help guide consideration of potential risks and benefits in the therapeutic decisionmaking process.

## Systematic Review of the Evidence

The following description of the SRE is an adaptation of the evidence report, including direct excerpts, submitted by the Blue Cross Blue Shield Association Evidence-Based Practice Center. (See Introduction, Methods.)

### I Methods of Literature Search

To be eligible for consideration in the SRE, each study was required to meet the following criteria:

- It reported on inhaled corticosteroid treatment.
- The treatment duration/observation was at least 1 year.
- For prospective studies:
  - Enrolled only patients younger than 18 years of age.
- OR
- Stratified outcomes for patients younger than 18 years of age and reported baseline demographics for the stratified subgroup.
- For retrospective studies:
  - Enrolled children and/or young adults younger than 40 years of age and indicated that a substantial proportion of the exposure to inhaled corticosteroids had been during childhood.
  - Study design was a comparative clinical trial, cohort study, case control study, or cross-sectional study.
- Reported on a group of at least 25 evaluable, similarly treated asthma patients per study arm.
- For growth outcomes:
  - Studies of short-term growth were restricted to randomized clinical trials.
  - Studies of long-term growth were restricted to studies that assessed final attained adult height and controlled for confounding variables.
  - For bone density, studies were restricted to controlled trials.
  - For subcapsular cataract, clinical series studies were also included.



- For hypothalamic-pituitary-adrenal (HPA) axis function, studies also were included that used a pre-post single-arm design, where baseline HPA axis function was measured before initiation of inhaled corticosteroids.

## I Summary of Findings

### Studies

The SRE addressed the long-term adverse effects of chronic inhaled corticosteroid use in children on four outcomes: vertical growth; bone mineral density; ocular toxicity, including posterior subcapsular cataract and glaucoma; and suppression of adrenal/pituitary axis. (See the key evidence tables in this section for a description of the studies reviewed for vertical growth [three retrospective cohort studies on final height]; bone-mineral density [two cross-sectional studies and one randomized controlled trial]; and HPA axis function [six studies, including three randomized controlled trials]). The difficulties of systematically assessing adverse effects are well known. Most clinical trials are not designed to specifically address adverse effects and thus may be statistically underpowered and of insufficient duration to detect long-term adverse effects. In addition, the results of this evidence review do not apply to adults. For the adult population, particularly elderly adults, adverse effects may differ qualitatively and quantitatively. For example, although effects on vertical growth are not a concern for adults, ocular toxicity is likely to occur more frequently as age increases.

### Results of Studies

The available evidence suggests that the use of inhaled corticosteroids at recommended doses does not have frequent, clinically significant, or irreversible effects on any of the outcomes reviewed. It is possible that chronic use of inhaled corticosteroids initiated in childhood and continued through adulthood might have cumulative effects that increase the relative risk of certain conditions—such as osteoporosis, cataracts, or glaucoma—in later life. However, none of the available studies had sufficient followup duration or numbers of patients to assess this possibility definitively. It is also likely that the probability of adverse effects is related to inhaled corticosteroids dosage. No studies identified in the published literature, however, were designed to test

the dose-response relationship of inhaled corticosteroids to adverse effects.

### Vertical Growth

The long-term prospective studies on growth involved budesonide, and the retrospective analyses included studies on beclomethasone, but the results have been generalized to all inhaled corticosteroid preparations. Although different preparations and delivery services may have a systemic effect at different doses, all short-term studies of numerous preparations suggest that the effect of inhaled corticosteroids on growth is a drug class effect.

Evidence addressing three measures of vertical growth in children was found: short-term growth velocity measured over a period of 1 year or less, growth velocity and change in height measured over longer duration (4 to 6 years), and final attained adult height. The evidence on short-term growth velocity is from a published meta-analysis, which pooled data from 5 randomized controlled trials representing 855 subjects, with a mean age of 9.5 years (Sharek and Bergman 2000). Evidence on growth velocity and height over a longer period of time is from the CAMP trial, comparing inhaled corticosteroids (budesonide), nedocromil, and placebo in 1,041 children with mild or moderate persistent asthma, who were followed for 4 to 6 years (CAMP 2000). For final attained adult height, evidence is from three retrospective cohort studies that adjusted for the potential confounding factor of parental height (Agertoft and Pedersen 2000; Silverstein et al. 1997; Van Bever et al. 1999). Together, these three studies included a total of 243 patients with asthma treated with inhaled corticosteroids, 154 asthmatic patients who had not been treated with inhaled corticosteroids, and 204 nonasthmatic controls.

Evidence on growth velocity when evaluated during the first year of therapy is consistent in showing a difference in height averaging approximately 1 cm between children treated with inhaled corticosteroids and controls. The magnitude of this change in height ( $\approx 0.5 \rightarrow 1.5$  cm) has varied between studies using different inhaled corticosteroid preparations, indicating that either the study design or specific steroid preparation/

dose may be important considerations (Doull et al. 1995; Allen et al. 1998; Verberne et al. 1997). In the only trial extending beyond 1 year (CAMP 2000), a difference consistent with this magnitude also occurred during the first year of the study. However, in subsequent long-term followup, the difference in growth velocity was not maintained; all groups had similar growth velocity at the end of treatment. At the end of the 4- to 6-year treatment period, there was still an approximately 1 cm difference in cumulative growth between the study groups, but a slight difference in bone age suggests the potential for catchup for the inhaled corticosteroid group.

The evidence on final adult height appears to be fairly consistent as well. However, this evidence is based on cohort studies that are subject to selection bias and the confounding effects of severity of asthma cannot be adjusted. Some comparisons in these studies also were limited by small sample size. Of the three studies, two showed no difference, and one showed a difference in final attained adult height between inhaled corticosteroid users and nonusers. However, the difference was much less than would be expected if a 1 cm/year growth velocity difference noted in the 1-year studies were maintained over several years.

#### Bone Mineral Density

The CAMP study followed children with mild or moderate persistent asthma and a mean age of approximately 9 years who were treated for 4 to 6 years with inhaled corticosteroids. This study, with large numbers, randomization, and assessment of longitudinal changes, provides strong evidence that there is no effect of inhaled corticosteroids on bone mineral density (BMD) in the doses given and in the duration in the study (CAMP 2002). One retrospective study of 30 young adults found a significant correlation between BMD and dose of inhaled corticosteroids among female patients (Ip et al. 1994). Such studies are subject to potential confounding because of unmeasured differences between groups that are risk factors for low BMD. In addition, the clinical significance of any observed differences in BMD are unknown. Subtle differences in BMD would not have a clinical impact until they were added to other risk factors such as

aging, and it is uncertain whether differences observed during young adulthood would persist into old age.

**Posterior Subcapsular Cataract and Glaucoma**  
Studies that report on the occurrence of posterior subcapsular cataracts consist mostly of small cohorts and cross-sectional studies (Allen et al. 1998; Tinkelman et al. 1993; Agertoft et al. 1998; Simons et al. 1993; Nassif et al. 1987; Abuekteish et al. 1995), with the exception of the CAMP study. The expected incidence rate of subcapsular cataract in any population of normal young children and adults is none. These studies are sufficient to rule out a large effect of inhaled corticosteroids on the short-term incidence of cataract, but they are not capable of detecting a small increase in risk of an event that has a baseline risk of essentially zero. In addition, several of the clinical trials that evaluated development of cataracts were of relatively short duration.

Two of these studies also reported on measurements of ocular pressure (Tinkelman et al. 1993; Nassif et al. 1987). The limited data available show no relationship between glaucoma or increased intraocular pressure and inhaled corticosteroids.

#### Effect on Hypothalamic-Pituitary-Adrenal Axis Function

Two types of evidence on the effects of inhaled corticosteroids on HPA axis function have been reported: three case reports of iatrogenic Cushing syndrome that were possibly related to inhaled corticosteroids (Zimmerman et al. 1998; Taylor et al. 1999; Priftis et al. 1991; Hollman and Allen 1988) and six controlled clinical trials regarding HPA axis function (Tinkelman et al. 1993; Nassif et al. 1987; Scott and Skoner 1999; Ribeiro 1993; Price et al. 1997; Gonzalez Perez-Yarza et al. 1996). Each study evaluated from one to three different measures of HPA axis function, with followup for at least 1 year after initiation of treatment.

The case reports show that systemic effects can occur in clinically detectable ways, with a strong case for causality indicated in the case studies by the accompanying laboratory tests and response when inhaled corticosteroids were withdrawn.

In the controlled clinical studies, four studies of serum control values identified no differences. However, three other studies used more sensitive tests of cortisol, such as 24-hour urinary cortisol, and two showed a statistically significant effect of inhaled corticosteroids. It should be noted that these statistically significant results occur as comparisons of mean values between groups. Few or no patients in most studies produce laboratory values out of the normal range. However, the clinical significance of these more sensitive indicators of adrenal function is unknown.

The results of the case reports appear to be causally attributable to inhaled corticosteroids based on clinical presentation, consistency with laboratory findings, and clinical response to reduction or withdrawal of treatment. Although the studies show that, on average, persons may only have clinically insignificant effects of inhaled corticosteroids on the HPA axis, some individuals may be acutely susceptible to their effects.

### Additional Literature/Information

Since the release of the EPR-2, a FDA-based committee convened to review the safety of inhaled corticosteroid therapy, with particular emphasis on growth effects. The FDA committee recommended inserting the following cautionary wording in package inserts for all (both nasal and oral) inhaled corticosteroid medications: "A reduction in growth velocity in children or teenagers may occur as a result of inadequate control of chronic diseases such as asthma or from use of corticosteroids for treatment. Physicians should follow closely the growth of adolescents taking corticosteroids by any route and weigh the benefits of corticosteroid therapy and asthma control against the possibility of growth suppression if an adolescent's growth appears slowed (<http://www.fda.gov>)."

Two additional studies on the effect of inhaled corticosteroids were completed after the SRE; the studies involved primarily adults but included some children and thus were considered by the Expert Panel. One report pertaining to the risk of cataract formation among patients 3 to 90 years of age was

based on a large retrospective cohort study in the United Kingdom-based General Practice Research Database population, with a nested case-control analysis among users of inhaled corticosteroids and patients without previous steroid use who were younger than 90 years of age. All users of inhaled corticosteroids were at a marginally increased risk of cataract formation (risk ratio = 1.3) compared to patients who did not use corticosteroids. Among individuals 40 years of age or older, the risk ratio increased as numbers of inhaled corticosteroid prescriptions increased after controlling for other variables. These trends were not evident for those individuals younger than 40 years of age (Jick et al. 2001).

A prospective cohort study on bone loss in women 18 to 45 years of age reported that bone-density loss at the total hip and the trochanter—but not at the femoral neck or spine—increased with the number of puffs per day of an inhaled corticosteroid (Israel et al. 2001). However, the clinical significance of these findings is uncertain because the rate of loss reported was small, any association of this small loss with increased risk of bone fracture has not been established, and the rates varied among the women taking the inhaled corticosteroids.

### Recommendations: EPR-2

Based on this information from the SRE and additional studies, the Expert Panel recommends the following text (the blue text is new text) as an update to pages 71 through 73 of EPR-2 (The Medications, Special Issues on Safety, Systemic Adverse Effects). This text updates—but does not change—the EPR-2 recommendations.

#### Linear Growth

A reduction in growth velocity in children or adolescents may occur as a result of inadequate control of chronic diseases such as asthma or from the use of corticosteroids for treatment. Overall, however, the available cumulative data in children suggest that, although low-to-medium doses of inhaled corticosteroids may have the potential of decreasing growth velocity, the effects are small, nonprogressive, and may be reversible (SRE-Evidence A, B, C).

The long-term prospective studies on growth involved budesonide, and the retrospective analyses included studies on beclomethasone, but the results have been generalized to include all inhaled corticosteroid preparations. Although different preparations and delivery devices may have a specific effect at different doses, the long-term effects on numerous preparations suggest that the effect of inhaled corticosteroids on growth is a drug-class effect. When high doses of inhaled corticosteroids are necessary to achieve satisfactory asthma control, the use of adjunctive long-term corticosteroid therapy is indicated in order to reduce the dose of inhaled corticosteroids and thus minimize possible dose-related effects on growth. Physicians should monitor the growth of children and adolescents receiving corticosteroids by accurate and weight-based benefits of corticosteroid therapy and asthma control against the possibility of growth suppression or if a child or an adolescent's growth appears slowed.

#### Bone Mineral Density

Low-to-medium doses of inhaled corticosteroids appear to have no serious adverse effects on BMD in children (SRE-Evidence A) (CAMP 2000). A small, dose-dependent reduction in BMD may be associated with inhaled corticosteroids in patients older than 18 years of age (SRE-Evidence C; Evidence B) (Ip et al. 1994; Israel et al. 2000), but the clinical significance of these findings is not clear.

#### Cataracts

In children, low-to-medium dose inhaled corticosteroid therapy has no significant effects on the incidence of subcapsular cataracts or glaucoma (SRE-Evidence A, C) (CAMP 2000; Jick et al. 2001). High (greater than 2000 mg) cumulative lifetime doses of inhaled corticosteroids may increase slightly the prevalence of cataracts as suggested in two retrospective studies of adult and elderly patients (SRE-Evidence C; Evidence C) (Cumming et al. 1997; Jick et al. 2001).

#### Hypothalamic-Pituitary-Adrenal Axis Function

The available evidence indicates that, on average, children may experience only clinically insignificant, if any, effects of low-to-medium dose inhaled corticosteroids on the HPA axis (SRE-Evidence A, C). Some individuals, however, may be more susceptible to their effects even at conventional doses.

#### Recommendations for Future Research

- What are the long-term effects of inhaled corticosteroid therapy on BMD and cataract formation if it is initiated at a young age and continued for prolonged periods of time?
- Are potential growth effects of inhaled corticosteroid therapy more pronounced during certain developmental periods (e.g., first 3 years of life, preadolescence)?

## Key Evidence Tables

Table 1-3. Differences in Adult Target Height in Cohort Studies

Study	Group (n) Comparison	Difference in (Adult Target) Height (cm) <sup>1</sup>
Silverstein, Yunginger, Reed et al. 1997	All asthmatics (n = 153) vs. nonasthmatics (n = 153)	0.2
	All corticosteroid users (n = 58) vs. noncorticosteroid asthmatics (n = 95)	-1.2
	Males: All corticosteroid users (n = 30) vs. noncorticosteroid asthmatics (n = 45)	-1.8
	Females: All corticosteroid users (n = 28) vs. noncorticosteroid asthmatics (n = 50)	-0.8
	Oral corticosteroid users (n = 40) vs. never used corticosteroids (n = 95)	-1.4
	Inhaled corticosteroid users (n = 18) vs. never used corticosteroids (n = 95)	-0.9
Van Bever, Desager, Lijssens et al. 1999	All inhaled corticosteroid users (n = 43) vs. never used corticosteroids (n = 42)	-2.54 <sup>2</sup>
	Males: Inhaled corticosteroid users (n = 23) vs. never used corticosteroids (n = 26)	-3.09 <sup>2</sup>
	Females: Inhaled corticosteroid users (n = 20) vs. never used corticosteroids (n = 16)	-1.99
Agertoft and Pedersen 2000	All inhaled corticosteroid users (n = 142) vs. noncorticosteroid using asthmatics (n = 18)	+0.5
	All inhaled corticosteroid users (n = 142) vs. healthy sibling control group (n = 51)	-0.6
	Males: All inhaled corticosteroid users (n = 86) vs. healthy sibling control group (n = 24)	-0.6
	Females: All inhaled corticosteroid users (n = 56) vs. healthy sibling control group (n = 27)	-0.8

<sup>1</sup> A negative number indicates that corticosteroid users had lower attained adult height than the comparison group, controlling for parental height.

<sup>2</sup> p < 0.05

## Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.



Table 1-4. Effects of Inhaled Corticosteroids on Bone Mineral Density

Citation	Treatment	Number Enrolled	Number Evaluable	Treatment Duration (years)	Bone	P Value	Comment
Agertoft, Larsen, and Pedersen 1998	Budesonide 504 mcg per day	157	157	3.0 (minimum)	Total body BMD: 0.92 g/cm <sup>2</sup>		No significant difference between groups or between boys and girls in bone mineral capacity or total bone calcium
	Nonsteroid asthma therapies	111	111	3.0 (minimum)	Total body BMD: 0.92 g/cm <sup>2</sup>	NS	Mean treatment time 4.4 (3–6) years
Ip, Lam, Yam, et al. 1994	Beclomethasone or budesonide	30	30	3.3	Spine: 0.944	0.041	Stratified by sex, all differences significant for females but not for males
					Femur Neck: 0.769	0.007	
					Trochanter: 0.676	0.034	
					Ward's Triangle: 0.729	0.016	
	Normal control subjects, matched by sex, age, BMI, menopausal status	30	30	NA	Spine: 1.011		
					Femur Neck: 0.835		
					Trochanter: 0.724		
					Ward's Triangle: 0.729		
Childhood Asthma Management Program Research Group 2000a	Budesonide 400 mcg/day	311	311	4–6	Change in spine BMD: 0.17 g/cm <sup>2</sup>	0.53 vs. placebo	
	Nedocromil 16 mg/day	312	312	4–6	Change in spine BMD: 0.17 g/cm <sup>2</sup>	0.15 vs. placebo	
	Placebo	418	418	4–6	Change in spine BMD: 0.18 g/cm <sup>2</sup>		

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.



Table 1-5. Effects of Inhaled Corticosteroids on HPA Function

Citation	Arms	Measure of HPA Axis Function	
Randomized Clinical Trials			
Scott and Skoner 1999	BUD 500 mcg/day (n = 132) vs. conventional treatment (n = 57)	Serum cortisol at baseline and 12 mo.  ACTH-stimulated cortisol at baseline and 12 mo.  Percentage of patients from normal to abnormal stimulation test between baseline and 12 mo.	
Price, Russell, Hindmarsh et al. 1997	FP 50 mcg/day (n = 36) vs. cromolyn 20 mg/day (n = 27)	Urinary cortisol geometric mean ratio between patient groups at 6 and 12 mo.	
Tinkelman, Reed, Nelson et al. 1993	BDP 34 mcg/day (n = 102) vs. theophylline (n = 93)	Serum cortisol at baseline, 6 and 12 mo.  ACTH-stimulated cortisol at baseline, 6 and 12 mo.	
Cross-Section Studies			
Gonzales Perez-Yarza, Mintegui, Garmendia et al. 1996	Budesonide or beclomethasone mean dose 676 +/- 280 mcg/day (range, 226-1800) (n = 250) vs. normal controls (n = 108)	Urinary cortisol  Number of abnormal ACTH stimulation tests in subset with urinary cortisols below 1 standard deviation	
Nassif, Weinberger, Sherman et al. 1987	Beclomethasone 358 mcg/day (n = 17) vs. Beclomethasone 726 mcg/day (n = 14) vs. asthmatic control group (n = 20) and normal control groups (n = 21)	Serum cortisol  Urinary cortisol	
Single Arm Pre-Post Study			
Ribiero 1993	Budesonide 200 mcg/day (n = 47)	Serum cortisol at baseline and 12 mo.  ACTH-stimulated cortisol at baseline and 12 mo.	

Results		P Value	Comments
BUD (0, 12 mo.): 320, 300 Conventional (0, 12 mo.): 250, 315	"No significant differences"	Subset of full trial	
BUD (0, 12 mo.): 695, 655 Conventional (0, 12 mo.): 690, 720	"No significant differences"	Subset of full trial	
BUD: 24% Conventional: 21%	"Not different"		
Ratio of urinary cortisol at 6 mo.: 0.85 Ratio of urinary cortisol at 12 mo.: 0.96	NS: 95% CI includes 1 NS: 95% CI includes 1		
BDP 336 mcg/day (0, 6, 12 mo.): 328, 306, 309 Theophylline (0, 6, 12 mo.): 309, 322, 334 BDP 336 mcg/day (baseline): 726 (6, 12 mo. NA) Theophylline (baseline): 723 (6, 12 mo. NA)	Not stated: "similar"  Not stated: "almost identical"		
BUD/BDP: 58.69 nmol/m <sub>2</sub> /day Control: 81.98 nmol/m <sub>2</sub> /day BUD/BDP group: 2 abnormal tests (3.1%) Control group: Not done	p <0.05  Not applicable	One of the two patients with abnormal test had chronic oral corticosteroids.	
BDP <450 mcg/day: 403 BDP >450 mcg/day: 353 Asthmatic controls: 353 Normal controls: 367  BDP <450 mcg/day: 22 mcg/g creatinine BDP >450 mcg/day: 16.5 mcg/g creatinine Asthmatic controls: 43 mcg/g creatinine Normal controls: 29.5 mcg/g creatinine	Not specifically stated: presumed NOT statistically significant  Text: "Statistically significant" from controls		
Basal cortisol (0, 12 mo.): 497, 497  4-hr. stimulated cortisol (0, 12 mo.): 1104, 1131 5-hr. stimulated cortisol (0, 12 mo.): 1242, 1380	Not stated, presumed not statistically significant  p = 0.02 for increase from baseline, both tests		

## Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

## References

- Abuekteish F, Kirkpatrick JN, Russell G. Posterior subcapsular cataract and inhaled corticosteroid therapy. *Thorax* 1995;50(6):674-6.
- Agertoft L, Larsen FE, Pedersen S. Posterior subcapsular cataracts, bruises and hoarseness in children with asthma receiving long-term treatment with inhaled budesonide. *Eur Respir J* 1998;12(1):130-5.
- Agertoft L, Pedersen S. Effect of long-term treatment with inhaled budesonide on adult height in children with asthma. *N Engl J Med* 2000;343(15):1064-9.
- Allen DB, Bronsky EA, LaForce CF, Nathan RA, Tinkelman DG, Vandewalker ML, Konig P. Growth in asthmatic children treated with fluticasone propionate. Fluticasone Propionate Asthma Study Group. *J Pediatr* 1998;132(3 Pt 1):472-7.
- Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.
- Childhood Asthma Management Program (CAMP) Research Group. Long-term effects of budesonide or nedocromil in children with asthma. *N Engl J Med* 2000;343(15):1054-63.
- Cumming RG, Mitchell P, Leeder SR. Use of inhaled corticosteroids and the risk of cataracts. *N Engl J Med* 1997;337(1):8-14.
- Doull IJ, Freezer NJ, Holgate ST. Growth of prepubertal children with mild asthma treated with inhaled beclomethasone dipropionate. *Am J Respir Crit Care Med* 1995;151(6):1715-19.
- Gonzalez Perez-Yarza E, Mitegui J, Garmendia A, Callen M, Reguilon MJ, Garrido A, Emparanza JI. The excretion of free cortisol in the urine in healthy children and in asthmatics treated with long-term inhaled glucocorticoids. *An Esp Pediatr* 1996;44(6):531-6.
- Hollman GA, Allen DB. Overt glucocorticoid excess due to inhaled corticosteroid therapy. *Pediatrics* 1988;81(3):452-5.
- Ip M, Lam K, Yam L, Kung A, Ng M. Decreased bone mineral density in premenopausal asthma patients receiving long-term inhaled steroids. *Chest* 1994;105(6):1722-7.
- Israel E, Banerjee TR, Fitzmaurice GM, Kotlov TV, LaHive K, LeBoff MS. Effects of inhaled glucocorticoids on bone density in premenopausal women. *N Engl J Med* 2001;345(13):941-7.
- Jick SS, Vasilakis-Scaramozza C, Maier WC. The risk of cataract among users of inhaled steroids. *Epidemiology* 2001;12(2):229-34.
- Nassif E, Weinberger M, Sherman B, Brown K. Extrapulmonary effects of maintenance corticosteroid therapy with alternate-day prednisone and inhaled beclomethasone in children with chronic asthma. *J Allergy Clin Immunol* 1987;80(4):518-29.

- Price JF, Russell G, Hindmarsh PC, Weller P, Heaf DP, Williams J. Growth during one year of treatment with fluticasone propionate or sodium cromoglycate in children with asthma. *Pediatr Pulmonol* 1997;24(3):178-86.
- Priftis K, Everard ML, Milner AD. Unexpected side-effects of inhaled steroids: a case report. *Eur J Pediatr* 1991;150(6):448-9.
- Ribeiro LB. Budesonide: safety and efficacy aspects of its long-term use in children. *Pediatr Allergy Immunol* 1993;4(2):73-8.
- Scott MB, Skoner DP. Short-term and long-term safety of budesonide inhalation suspension in infants and young children with persistent asthma. *J Allergy Clin Immunol* 1999;104(4 Pt 2):200-9.
- Sharek PJ, Bergman DA. The effect of inhaled steroids on the linear growth of children with asthma: a meta-analysis. *Pediatrics* 2000;106(1):E8.
- Silverstein MD, Yunginger JW, Reed CE, Petterson T, Zimmerman D, Li JT, O'Fallon WM. Attained adult height after childhood asthma: effect of glucocorticoid therapy. *J Allergy Clin Immunol* 1997;99(4):466-74.
- Simons FE, Persaud MP, Gillespie CA, Cheang M, Shuckett EP. Absence of posterior subcapsular cataracts in young patients treated with inhaled glucocorticoids. *Lancet* 1993;342(8874):776-8.
- Taylor DA, Jensen MW, Kanabar V, Engelstatter R, Steinijans VW, Barnes PJ, O'Connor BJ. A dose-dependent effect of the novel inhaled corticosteroid ciclesonide on airway responsiveness to adenosine-5'-monophosphate in asthmatic patients. *Am J Respir Crit Care Med* 1999;160(1):237-43.
- Tinkelman DG, Reed CE, Nelson HS, Offord KP. Aerosol beclomethasone dipropionate compared with theophylline as primary treatment of chronic, mild to moderately severe asthma in children. *Pediatrics* 1993;92(1):64-77.
- Van Bever HP, Desager KN, Lijssens N, Weyler JJ, Du Caju MV. Does treatment of asthmatic children with inhaled corticosteroids affect their adult height? *Pediatr Pulmonol* 1999;27(6):369-75.
- Verberne AA, Frost C, Roorda RJ, van der Laag H, Kerrebijn KF. One year treatment with salmeterol compared with beclomethasone in children with asthma. The Dutch Paediatric Asthma Study Group. *Am J Respir Crit Care Med* 1997(3 Pt 1):156:688-95.
- Zimmerman B, Gold M, Wherrett D, Hanna AK. Adrenal suppression in two patients with asthma treated with low doses of the inhaled steroid fluticasone propionate. *J Allergy Clin Immunol* 1998;101(3):425-6.

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## Combination Therapy: Addition of Other Long-Term-Control Medications to Inhaled Corticosteroids

### Question

*In patients with moderate persistent asthma who are receiving inhaled corticosteroids, does addition of another long-term-control agent improve outcomes?*

### Summary Answer: Question

Strong evidence consistently indicates that long-acting inhaled  $\beta_2$ -agonists added to low-to-medium-dose inhaled corticosteroids improve outcomes (SRE-Evidence A). Adding a leukotriene modifier or theophylline to inhaled corticosteroids or doubling the dose of inhaled corticosteroids also improves outcomes, but the evidence is not as substantial (SRE-Evidence B). The EPR-2 recommendations for moderate persistent asthma have been revised: The preferred treatment for adults and children older than 5 years of age is the addition of long-acting inhaled  $\beta_2$ -agonists to low-to-medium-doses of inhaled corticosteroids. Adjunctive therapy combinations have not been studied in children younger than 5 years of age. For this age group, it is the opinion of the Expert Panel that there are two preferred options for treating moderate asthma: either the addition of long-acting inhaled  $\beta_2$ -agonists to a low dose of inhaled corticosteroids or medium-dose inhaled corticosteroids as monotherapy.

### Rationale for the Question

There are an increased number of studies evaluating combination therapy primarily as a result of the development of fixed-dose combinations of the long-acting inhaled  $\beta_2$ -agonists and inhaled corticosteroids (salmeterol plus fluticasone propionate, now FDA-approved, and formoterol plus budesonide, under development). The ongoing preference to minimize the dose of corticosteroids, especially for patients taking high doses, and to reduce the possibility of adverse side effects, has stimulated studies of adjunctive therapies. The question

of interest is whether, for patients requiring more than low doses of inhaled corticosteroids, equal or better asthma control could be achieved by adding an additional medication rather than by increasing the dose of inhaled corticosteroids. An extensive body of literature addressing the question of adjunctive therapy has become available since the publication of EPR-2 and has thus warranted Expert Panel Review.

### Systematic Review of the Evidence

The following description of the SRE is an adaptation of the evidence report, including direct excerpts, submitted by the Blue Cross Blue Shield Association Evidence-Based Practice Center. (See Introduction, Methods.)

#### Methods of Literature Search

The SRE divided the studies into three study design categories:

1. The addition of a long-term-control medication to a fixed dose of inhaled corticosteroids compared with the same dose of inhaled corticosteroids alone. This design simply assesses whether combination therapy is better than monotherapy with inhaled corticosteroids. The potential bias from this study design is seen when patients can be controlled on inhaled corticosteroids alone, resulting in a negative study because of the inability to improve.
2. The addition of a long-term-control medication to inhaled corticosteroids with subsequent downward titration of the dose of inhaled corticosteroids to the lowest dose that maintains control. This design is even more problematic because it may be raising a fundamentally different question—i.e., “Can the other long-term-control medication act as a substitute for the inhaled corticosteroids following initial control of the asthma?” However, if the goal is simply to lower the dose of inhaled corticosteroids by some increment (usually half), then the study design addresses the primary question more directly.



3. The addition of the long-term-control medication compared with increasing the dose of inhaled corticosteroids to improve asthma control. This design most directly addresses the question, because eligible patients first demonstrated a lack of adequate control during an open run-in period on inhaled corticosteroids. The definition of inadequate control varied among studies, however, and this variance could introduce some bias.

In addition to the eligibility criteria for selecting studies related to all topics in the SRE (described in the Introduction), the criteria for selecting studies for this question were as follows:

- Study comparisons included:
  - Inhaled corticosteroids alone compared to inhaled corticosteroids plus leukotriene modifiers, or long-acting beta<sub>2</sub>-agonists, or theophylline
- OR
- Two different long-term-control medications in patients using inhaled corticosteroids
- OR
- The addition of an alternative medication to an increased dose of inhaled corticosteroids for patients already on inhaled corticosteroids.
- Treatment duration was at least 4 weeks.
- At least 90 percent of patients in the study were on inhaled corticosteroids, or the subgroup of patients on inhaled corticosteroids was analyzed separately, and this subgroup otherwise met the eligibility criteria for this question.
- No more than 10 percent of the patients in the population or in a subgroup were on oral corticosteroids.

## I Summary of Findings

### Studies

The majority of the studies reviewed by the SRE fit into study design categories 1 and 3. Thirty-nine studies involving 45 comparisons and a total of 9,020 patients were selected for the SRE. (See the key evidence tables in this section.) Overall, 34 of the 45 comparisons evaluated the addition of a

long-acting beta<sub>2</sub>-agonist to inhaled corticosteroids. All but one of the studies were randomized trials. The following comparisons were made:

- Twenty-six compared the addition of a drug to a fixed dose of inhaled corticosteroids (18 [3,163 patients] compared long-acting inhaled beta<sub>2</sub>-agonists; 4 [234 patients] compared theophylline; and 4 [885 patients] compared LTRAs).
- Four compared a titrated dose of inhaled corticosteroids after the addition of a drug (3 [268 patients] compared long-acting inhaled beta<sub>2</sub>-agonists; 1 [226 patients] compared LTRA).
- Fifteen compared a low-to-moderate dose of inhaled corticosteroids with an additional drug to high-dose inhaled corticosteroids (13 [4,285 patients] compared long-acting inhaled beta<sub>2</sub>-agonists and 2 [252 patients] compared theophylline).
- No studies were found that compared long-acting oral beta<sub>2</sub>-agonists.
- No studies meeting SRE quality criteria were found that compared the addition of cromolyn or nedocromil.

### Results of Studies

**Addition of long-acting inhaled beta<sub>2</sub>-agonists**  
A sufficient number of quality studies in both design categories 1 and 3 were completed to enable meta-analyses of lung function and as-needed short-acting beta<sub>2</sub>-agonist use outcomes in each category. (See the key evidence tables in this section for a description of eligible studies.) Both the systematic review and meta-analyses confirmed the superiority of combination therapy to inhaled corticosteroids monotherapy. In particular, the findings of the meta-analysis for the addition of long-acting inhaled beta<sub>2</sub>-agonist compared with increasing the inhaled corticosteroid dosage were consistent with a previously reported meta-analysis (Shrewsbury et al. 2000). In addition to similar findings on lung function, Shrewsbury and colleagues had access to the original data and were able to assess the rate of asthma exacerbations, reporting a positive benefit of the combination

therapy. The data are robust and convincing that the addition of long-acting inhaled  $\beta_2$ -agonists to inhaled corticosteroids improves lung function and asthma control in patients inadequately controlled with low-to-medium doses of inhaled corticosteroids.

Of note is the paucity of pediatric trials in the database. One pediatric study by Verberne et al. (1998) was completed in older children (mean 11 years of age). Following a 6-week run-in, 120 patients were randomized to either low-dose inhaled corticosteroid—beclomethasone dipropionate (BDP) (400 mcg/day), medium-dose BDP (800 mcg/day), or low-dose BDP plus the long-acting inhaled  $\beta_2$ -agonist salmeterol for 1 year. No significant difference was found among any of the three arms in postbronchodilator FEV<sub>1</sub> or PC20 FEV<sub>1</sub>, methacholine provocation. These results suggest that the children's asthma was adequately controlled with low-dose inhaled corticosteroids and that the addition of the long-acting inhaled  $\beta_2$ -agonist neither improved nor worsened airway responsiveness. Thus, due to the design, this study cannot refute the potential benefit of the drug combination for those children inadequately controlled on low-dose inhaled corticosteroids alone.

A multicenter double-blind trial of salmeterol as added therapy for children who were not well controlled with inhaled corticosteroids (mean dose of 750 mcg/day) demonstrated significant improvement in morning PEF and symptom-free days in the long-acting inhaled  $\beta_2$ -agonist plus inhaled corticosteroid group, compared to the placebo plus inhaled corticosteroid group (Russell 1995). Although this study did not compare the addition of a long-acting inhaled  $\beta_2$ -agonist to an increased dose of inhaled corticosteroids, the patients were already receiving doses of inhaled corticosteroids ranging from 400 to 2,400 mcg a day. Thus, this study established a need for further asthma control in children already receiving inhaled corticosteroids; it also more directly addresses the question posed by the SRE.

**Addition of long-acting oral  $\beta_2$ -agonists**  
No studies were found.

**Addition of cromolyn/nedocromil**

No studies meeting the quality criteria of the SRE were found. No new studies since the publication of the EPR-2 were found.

**Addition of theophylline**

Six studies evaluated the addition of theophylline, including two more recent studies that compared the addition to increased inhaled corticosteroid dosage. The results indicate that the combination of drugs and the increased dose of the inhaled corticosteroids result in equivalent outcomes, suggesting that theophylline has only a modest steroid-sparing effect. None of the four studies (two in children 6 to 19 years of age) comparing the addition of theophylline to a fixed dose of inhaled corticosteroids met the quality criteria of the SRE, because all had study-design and statistical problems. No studies were found that included children younger than 6 years of age.

**Addition of leukotriene modifiers**

Five published studies evaluated the addition of leukotriene modifiers to fixed doses of inhaled corticosteroids; none compared the combination to increasing the dose of inhaled corticosteroids. Two of these studies used pranlukast, an LTRA unavailable in the United States, and one used zafirlukast in a dose four times the dosage recommended on the package label. None of the studies included children younger than 12 years of age. The most relevant of the five studies (Laviolette et al. 1999), which contributed the most patients and had the longest duration, failed to meet the definition of high quality for the SRE because it met only one of the quality indicators (double blinding). Limitations of these studies preclude definitive conclusions, but they reveal a trend showing improvement in lung function and, in some, symptoms from the combination of leukotriene modifiers and inhaled corticosteroids compared with a fixed dose of inhaled corticosteroids alone.

**Addition of an adjunctive agent and down titration of the inhaled corticosteroids**

This group of studies is discussed separately, as some of the trials were designed to ask a fundamentally different question (i.e., could the adjunctive therapy ultimately replace inhaled

corticosteroid therapy?). An example is the study that attempted to wean patients from the inhaled corticosteroids after beginning a long-acting inhaled beta<sub>2</sub>-agonist until they had an exacerbation or the inhaled corticosteroid therapy was discontinued (McIvor et al. 1998). Ten of the 13 patients in the long-acting inhaled beta<sub>2</sub>-agonist arm experienced an exacerbation only after discontinuing their inhaled corticosteroids, providing further evidence that the long-acting inhaled beta<sub>2</sub>-agonist should not be used as a substitute for anti-inflammatory therapy. One trial attempted to wean patients from the inhaled corticosteroids after addition of the LTRA montelukast, with the goal of maintaining adequate asthma control (Lofdahl et al. 1999). The mean percentage reduction in the dose of inhaled corticosteroids was 47 percent—a 17 percent increase over placebo—and 40 percent of patients were able to discontinue their inhaled corticosteroids compared with 29 percent in the placebo arm, which was not statistically significant. Thus, data are inconclusive about the “steroid sparing” effect of adjunctive therapy, and data show that patients cannot be entirely weaned from inhaled corticosteroids. In addition, data from these studies are insufficient to determine the relative “steroid-sparing” effect of the various adjunctive therapies. Finally, none of the studies included children younger than 5 years of age.

### Additional Literature/Information

In addition to reviewing studies published after the SRE, the Expert Panel considered four other issues relevant to the question of the use of combination therapy for the treatment of persistent asthma: the effect of the different combinations on the rate of exacerbations of asthma; the comparison of different combinations to determine relative effectiveness; the use of combination therapy in children 5 years of age and younger; and the use of combination therapy in severe persistent asthma.

#### Studies Published After the SRE

The addition of montelukast to inhaled corticosteroids was evaluated in 279 children 6 to 14 years of age with moderate asthma whose symptoms were not completely controlled on 400 mcg

budesonide daily (Simons et al. 2001). This study was a double-blinded, randomized, placebo-controlled, crossover trial with a 4-week open-label run-in period to establish the need for adjunctive therapy. Each treatment period also consisted of 4 weeks. The trial had sufficient power (95 percent) to detect a 4.4-percent difference between the placebo and the active drug in the primary end point, FEV<sub>1</sub> percent predicted. In the intention-to-treat analysis, no significant difference was found between the placebo and montelukast for the primary end point (1.3 percent difference). A post hoc analysis of the data revealed a statistically significant 1.9 percent difference between the active drug and the placebo. Other significant differences reported in favor of montelukast were a decrease in beta<sub>2</sub>-agonist usage (.33 puffs/day difference) and exacerbation days that also were defined by beta<sub>2</sub>-agonist usage—an improvement in morning and evening PEFs (9.7 L/min and 10.7 L/min, respectively). It was not indicated whether these were intention-to-treat analyses. Outcomes found to be the same at the end of the study included worsening asthma, global evaluations, number of asthma attacks requiring intervention, and quality of life.

Another study compared the addition of theophylline to low-dose BDP (400 mcg daily) with increasing the dose of BDP to 1,000 mcg daily or maintaining patients on the low-dose BDP alone for 7 months (Lim et al. 2000). The study found no difference between the high-dose inhaled corticosteroids and the theophylline group for any outcome, thus confirming the SRE findings.

#### Effect of Combination Therapy on the Rate of Exacerbations of Asthma

Reduction in the rate of asthma exacerbations has been suggested as a surrogate for an anti-inflammatory effect. Compared with placebos, leukotriene modifiers have been reported to reduce the number of exacerbations treated with prednisone (zileuton, zafirlukast, and montelukast package inserts). Both of the long-acting inhaled beta<sub>2</sub>-agonists—formoterol and salmeterol—have been reported to reduce exacerbations of asthma when administered in conjunction with inhaled corticosteroids (Pauwels et al. 1997; Shrewsbury et al.

2000). In one study, the addition of formoterol to either low-dose (100 mcg bid) or high-dose (400 mcg bid) budesonide significantly reduced both mild and severe exacerbations. Further, fewer exacerbations occurred in the high-dose inhaled corticosteroid group compared with the lower dose group, though statistical analysis was not done (Pauwels et al. 1997). A meta-analysis of studies in which the addition of salmeterol to a lower dose of inhaled corticosteroids was compared with a higher dose of inhaled corticosteroids demonstrated that exacerbations were significantly lower with the combination therapy (Shrewsbury et al. 2000).

It has been suggested that this reduction in exacerbations may be attributed to an enhanced corticosteroid effect due to priming of the glucocorticoid receptor by the long-acting inhaled beta<sub>2</sub>-agonist (Eickelberg et al. 1999). Two recently published studies (Lazarus et al. 2001; Lemanske et al. 2001) also are pertinent to the issue of using asthma exacerbation as an outcome. In the first trial, those patients adequately controlled on low-dose inhaled corticosteroids were left on the inhaled corticosteroids, switched to the long-acting beta<sub>2</sub>-agonist salmeterol, or switched to placebo. Although the conventional outcomes (morning and evening PEFs) for the salmeterol and inhaled corticosteroid arms were not different, the salmeterol group had a significantly greater number of exacerbations and treatment failures—again demonstrating that the long-acting inhaled beta<sub>2</sub>-agonists cannot substitute for inhaled corticosteroids (Lazarus et al. 2001). The companion study evaluated the ability to reduce the dose of inhaled corticosteroids following the introduction of a long-acting inhaled beta<sub>2</sub>-agonist in those patients initially suboptimally controlled on the inhaled corticosteroids (Lemanske et al. 2001). In this group, the dose of inhaled corticosteroids was reduced by one-half in those patients responding to the addition without any significant change in asthma control, yet a significant treatment failure rate was noted when the inhaled corticosteroids were stopped.

Although clinical studies in the SRE suggest that the addition of a long-acting inhaled beta<sub>2</sub>-agonist to a low-to-medium dose of inhaled corticosteroids

is the most effective treatment for moderate persistent asthma (step 3 care), there may be situations where both the addition of a long-acting inhaled beta<sub>2</sub>-agonist and an increase in the dose of inhaled corticosteroids are indicated. The studies of Sont et al. (1999) and Pauwels et al. (1997) support the added benefit of a higher dose of inhaled corticosteroids in reducing asthma exacerbations. Thus, for patients considered to be at higher risk for exacerbations (suggested by a history of repeated short courses of prednisone, emergency department visits, or hospitalizations), both the addition of a long-acting inhaled beta<sub>2</sub>-agonist and an increase in the dose of inhaled corticosteroids may be indicated.

#### Comparison of Combinations To Determine Relative Effectiveness

Not included in the SRE were direct comparative studies of the effectiveness of the various drugs used as adjuncts to inhaled corticosteroids. Studies comparing the long-acting inhaled beta<sub>2</sub>-agonist to sustained-release theophylline are numerous (Davies et al. 1998), and generally involve patients receiving inhaled corticosteroids. A meta-analysis of these studies (Davies et al. 1998) demonstrated that both pulmonary function and asthma symptoms showed more improvement with the long-acting inhaled beta<sub>2</sub>-agonist as adjunctive therapy than with theophylline. In the three published studies included in the meta-analysis, between 50 percent and 97 percent of the subjects were receiving regular inhaled corticosteroid therapy (Fjellbirkeland et al. 1994; Muir et al. 1992; Paggiaro et al. 1996).

A comparison of the addition of the long-acting beta<sub>2</sub>-agonist salmeterol to the addition of the LTRA zafirlukast (Busse et al. 1999) also examined a mixed population; however, this study was not included in the SRE because more than 80 percent of the patients in both arms were using inhaled corticosteroids, rather than 90 percent required by the SRE selection criteria. The study otherwise met the criteria for a high-quality study and should be considered. The results indicate that salmeterol improved both pulmonary function and asthma symptoms significantly more than did zafirlukast.



Another direct comparison of long-acting inhaled  $\beta_2$ -agonists and a leukotriene modifier as combination therapy was published after the SRE (Nelson et al. 2000). This study also met the SRE criteria for high quality and should be considered. The investigators evaluated patients who were still symptomatic on low-dose inhaled corticosteroids (fluticasone 88 mcg bid), before and after the addition of the long-acting  $\beta_2$ -agonist salmeterol or the LTRA montelukast over 3 months. Those patients receiving salmeterol plus fluticasone, compared with those on montelukast and fluticasone, had greater improvement in pulmonary function and in some asthma symptoms, and experienced significantly fewer exacerbations.

Although the addition of sustained-release theophylline or a leukotriene modifier to treatment with inhaled corticosteroids generally is not as effective as the addition of a long-acting inhaled  $\beta_2$ -agonist, there may be circumstances when these combinations would be indicated for selected patients. Among the considerations favoring one of these alternative combinations would be the patient's intolerance of the side effects of the long-acting inhaled  $\beta_2$ -agonist, marked preference for oral therapy, demonstration of superior responsiveness to the alternate class of drug, as well as financial considerations (theophylline is the least expensive). Finally, although the recently marketed fixed-dose combination of fluticasone propionate and salmeterol in a DPI may provide an advantage in terms of ease of use (one inhaler instead of two), there is no evidence of superiority of this particular combination over that of other inhaled corticosteroids and long-acting inhaled  $\beta_2$ -agonists.

#### Combination Therapy in Children 5 Years of Age and Younger

None of the adjunctive therapy combinations have been adequately studied in children 5 years of age and younger. Indeed, only one study, a study adding the long-acting inhaled  $\beta_2$ -agonist salmeterol to inhaled corticosteroids, included patients as young as 4 years of age (Russell 1995). The lower age limit of all other combination therapy studies in children is 6 years of age (Simons et al. 2001; Meltzer et al. 1992; Nassif

et al. 1981). The data are thus inadequate to provide definitive recommendations on combination therapy in young children, and recommendations must be extrapolated from studies in older children and adults, which support the combination of inhaled corticosteroids and long-acting inhaled  $\beta_2$ -agonists. Because patients in this age range may be at greater risk for systemic effects from high doses of inhaled corticosteroids, the use of combination therapy seems prudent when goals of therapy are not attained with low or the lower range of medium doses of inhaled corticosteroids. However, as noted in the section on effectiveness of long-term-control medications, there are no data available on the use of long-acting inhaled  $\beta_2$ -agonists in infants and young children, whereas studies of medium doses of inhaled corticosteroids demonstrate effectiveness in this age group.

The following medications have been FDA-approved for young children: the inhaled corticosteroids budesonide nebulizer solution approved for children 1 to 8 years of age and fluticasone DPI approved for children 4 years of age and older; the long-acting inhaled  $\beta_2$ -agonist salmeterol DPI approved for children 4 years of age and older; and, based on safety data rather than efficacy data, the LTRA montelukast 4 mg chewable approved for children 2 to 6 years of age.

#### Combination Therapy in Patients With Severe Persistent Asthma

Current recommendations for treatment include adding oral systemic corticosteroids if a patient cannot achieve and maintain control with high doses of inhaled corticosteroids and long-acting bronchodilators. An alternative approach may be to add a third long-term-control medication to a combination of medium-to-high-dose corticosteroids and long-acting inhaled  $\beta_2$ -agonists in severe persistent asthma. However, few trials regarding this approach and of sufficient quality are available. A double-blind, crossover trial of LTRA (10 mg montelukast or placebo) in 72 adults with severe persistent asthma found no benefit from the addition of montelukast to other medication (Robinson et al. 2001). In this study, the concurrent medication varied among the patients: All patients received medium-to-high-dose inhaled

corticosteroids; 85 percent also received either theophylline, a long-acting inhaled  $\beta_2$ -agonist, or both; and 47 percent also received oral systemic corticosteroids. No attempt was made to eliminate the oral corticosteroids. The treatment period of 14 days for LTRA and 14 days for placebo was relatively short, although leukotriene modifiers usually produce a rapid response. This study indicates that there is no additional benefit to adding LTRA as a third medication. Similar controlled clinical trials have not been conducted to evaluate other long-term-control medications added to the combination of medium-to-high doses of inhaled corticosteroids and long-acting inhaled  $\beta_2$ -agonists in severe persistent asthma. Until more research is conducted, recommendations for managing severe persistent asthma are based on extrapolations from studies of the combination of inhaled corticosteroids and one other long-term-control medication in treating moderate persistent asthma.

### Recommendations for EPR update

Based upon the assessment of evidence provided by the SRE and the additional evidence considered by the Expert Panel, the following changes to step 3 care in EPR-2 are recommended:

- The preferred treatment for those adults and children older than 5 years of age whose asthma is inadequately controlled on low-dose inhaled corticosteroids is combination therapy: the addition of a long-acting inhaled  $\beta_2$ -agonist (SRE-Evidence A) to a low-to-medium dose of inhaled corticosteroids. Scientific evidence from studies of children older than 12 years of age and adults indicates that patients with moderate persistent asthma benefit from two different types of daily medication in order to achieve and maintain optimal control of their asthma: (1) medication aimed at suppressing underlying airway inflammation and (2) a medication whose primary action is bronchodilation. This approach is preferred to increasing the dose of inhaled corticosteroids.

The exception is indicated for those patients who experience recurring severe exacerbations that

require oral prednisone, emergency department visits, or hospitalizations. For these patients, increasing the dose of inhaled corticosteroids along with the addition of a long-acting inhaled  $\beta_2$ -agonist should be considered (SRE-Evidence B).

For children 5 years of age or younger, combination therapy has not been adequately studied. Therefore, recommendations for step 3 care for this age group are based on extrapolations of data from older children and adults, as well as expert opinion. For children 5 years of age and younger with moderate persistent asthma, there are two equally preferred options: low-dose inhaled corticosteroids and a long-acting  $\beta_2$ -agonist (Evidence B, extrapolation from studies in older children and adults) OR inhaled corticosteroids as monotherapy with an increase of the dose within the medium-dose range (Evidence D).

- Alternative—but not preferred—approaches that may be considered include doubling the dose of inhaled corticosteroids within the medium-dose range (this is an alternative but not preferred option for older children and adults; for children 5 years of age and younger, increasing the inhaled corticosteroid dose is an equally preferred option); adding sustained-release theophylline; or adding a leukotriene modifier (SRE-Evidence B). Leukotriene modifiers or theophylline may be considered if the patient displays intolerance of long-acting inhaled  $\beta_2$ -agonists, has a marked preference for oral therapy, and demonstrates superior responsiveness to the alternative class of drug through a therapeutic trial. Other issues may include financial considerations (theophylline is the least expensive).
- The recommendations for the use of nedocromil and long-acting oral  $\beta_2$ -agonists as alternatives to increasing the dose of inhaled corticosteroids are untenable at this time due to lack of data and should be removed as therapeutic options.

Specifically, the Expert Panel recommends that step 3 in figure 3–4b, Stepwise Approach for Managing Asthma in Adults and Children Older Than 5 Years of Age, be revised as follows with the revision noted in blue text.



Figure 3–6. Stepwise Approach for Managing Infants and Young Children (5 Years of Age and Younger) With Acute or Chronic Asthma. (See Medications: Effectiveness in Children on page 25 of this report for revisions to step 3.)

Figure 3–4b. Stepwise Approach for Managing Asthma in Adults and Children Older Than 5 Years of Age: Treatment (pages 84 through 85 in EPR-2)

### Step 3: Moderate Persistent

#### Daily Medication:

##### Preferred treatment:

Low-to-medium-dose inhaled corticosteroids and long-acting inhaled beta<sub>2</sub>-agonists

##### Alternative treatment (listed alphabetically):

Increase inhaled corticosteroids within medium-dose range

OR

Low-to-medium-dose inhaled corticosteroids and either a leukotriene modifier OR theophylline

If needed (particularly in patients with recurring severe exacerbations)

##### Preferred treatment:

Increase inhaled corticosteroids within medium-dose range and add a long-acting beta<sub>2</sub>-agonist

##### Alternative treatment:

Increase inhaled corticosteroids within medium-dose range and add either a leukotriene modifier

OR

Theophylline

### Step 4: Severe Persistent

#### Daily Medication:

##### Preferred treatment:

High-dose inhaled corticosteroids

AND

Long-acting inhaled beta<sub>2</sub>-agonists

AND, if needed

Corticosteroid tablets or syrup long term (1 to 2 mg/kg/day; generally do not exceed 60 mg/day). (Make repeat attempts to reduce systemic corticosteroids and maintain control with high-dose inhaled corticosteroids.)

The text in EPR-2 on pages 93 and 94 regarding step 3 and step 4 care for adults and children older than 5 years of age should be revised as follows, with the blue text indicating new text. (See Medications: Effectiveness in Children on page 25 for revisions to step 3 for children 5 years of age and younger.)

### Step 3: Moderate Persistent Asthma

Consultation with an asthma specialist may be considered because the therapeutic options at this juncture pose a number of challenging risk-benefit outcomes. Before increasing therapy, however, the clinician should review the patient's inhaler technique and adherence, as well as determine whether environmental factors are contributing to the patient's worsening asthma. If a step-up in therapy is required, there are at least four options for initiating therapy.

One option is to add a long-acting inhaled beta<sub>2</sub>-agonist to a low-to-medium-dose of inhaled corticosteroids (see Evidence Table B). This is the preferred treatment. Early investigations suggested that the addition of a long-acting inhaled beta<sub>2</sub>-agonist to a low-to-medium-dose of inhaled corticosteroids (Green et al. 1994) or medium-dose of inhaled corticosteroids (Shrewsbury et al. 1996) resulted in greater improvement in lung function and overall control than doubling the dose of inhaled corticosteroids. Since that time, numerous studies have confirmed the superiority of combination therapy over increasing the dose of inhaled corticosteroids, even for reducing severe asthma exacerbations (SRE 2004; Shrewsbury et al. 2000). Use of combination therapy has not been shown to mask worsening of inflammation and asthma. Indeed, the combination has consistently been shown to reduce the number of severe asthma exacerbations (Pauwels et al. 1997; Shrewsbury et al. 2000). This approach has proved so successful that it has spawned the development of two fixed-dose combinations of long-acting inhaled beta<sub>2</sub>-agonists and inhaled corticosteroids in one inhaler, one currently marketed. The fixed-dose combination may be easier to use and hence facilitate adherence to the regimen, but there is no evidence of clinical superiority over using the inhaled corticosteroids and long-acting inhaled beta<sub>2</sub>-agonists in separate inhalers.

OR

- Increase the dose of inhaled corticosteroids *and* add a long-acting inhaled beta<sub>2</sub>-agonist (SRE-Evidence B). This approach should be reserved for those patients experiencing recurring severe exacerbations, frequent oral prednisone, emergency department visits, or hospitalizations. In a 1-year combination therapy, the addition of a long-acting inhaled beta<sub>2</sub>-agonist to either low-dose or medium-dose inhaled corticosteroids significantly reduced the number of mild and severe exacerbations (Pauw et al. 1997). In addition, fewer patients in the high-dose inhaled corticosteroid group compared with the lower-dose group, although this analysis was not done.

OR

- Give inhaled corticosteroids alone or in combination by increasing the dose within the medium-dose range (SRE-Evidence B). This approach is another preferred treatment option for children; it is an alternative, but not preferred, treatment option for older children and adults. Studies of adults in which the dose of inhaled corticosteroids was at least doubled consistently demonstrate improved lung function and other outcomes in those patients not completely controlled on low-to-medium-doses of inhaled corticosteroids, but these results are consistently less effective than adding a long-acting inhaled beta<sub>2</sub>-agonist (SRE-Evidence A, B).

OR

- Add a leukotriene modifier or theophylline to inhaled corticosteroids (SRE-Evidence B; Evidence B). The addition of leukotriene modifiers and theophylline has produced modest improvement in lung function and some other outcomes in patients not completely controlled on inhaled corticosteroids. The addition of theophylline, however, has not been shown to be more effective than doubling the dose of inhaled corticosteroids (Evans et al. 1997; Ukena et al. 1997). The leukotriene modifiers have produced improvements in lung function and in some but not all measures of asthma control in patients incompletely controlled on inhaled corticosteroids (Laviolette et al. 1999). In addition, the leukotriene modifiers allow slightly more patients

to be taken off inhaled corticosteroids than does placebo (11 percent difference) (Lofdahl et al. 1999). The addition of the leukotriene modifiers to inhaled corticosteroids has not been compared with doubling the dose of inhaled corticosteroids. Direct comparisons of the addition of a leukotriene modifier or a long-acting inhaled beta<sub>2</sub>-agonist to therapy for patients incompletely controlled on inhaled corticosteroids show significantly greater improvement in lung function and other measures of asthma control for patients receiving the long-acting inhaled beta<sub>2</sub>-agonist and inhaled corticosteroid combination (Busse et al. 1999; Nelson et al. 2000). Thus, although the combination of inhaled corticosteroids and either theophylline or leukotriene modifier is not the preferred approach, considerations favoring one of these alternative combinations would be the patient's intolerance of the side effects of the long-acting inhaled beta<sub>2</sub>-agonist, marked preference for combination therapy, and demonstration of superior responsiveness to the alternative class of drug, as well as financial considerations (theophylline is more expensive).

Issues for children. Recommendations on combination therapy for children younger than 12 years of age with moderate persistent asthma are based on extrapolations from studies in older children and adults and on expert opinion (Evidence B, D). No combination therapy combinations have been adequately studied in children younger than 6 years of age and they have not been studied at all in children younger than 4 years of age. The negative results of combination therapy in children with mild or moderate persistent asthma failed to establish a need in the study participants at baseline for more therapy than low-dose inhaled corticosteroids and thus did not sufficiently address the question of combination therapy (Verberne et al. 1998). In one study in children 4 to 16 years of age with moderate or severe asthma, the addition of a long-acting beta<sub>2</sub>-agonist produced a clear benefit compared to placebo (Russell et al. 1995). In a recent crossover comparison of children 6 to 14 years of age on inhaled corticosteroids, no significant difference was found with the addition of the LTRA montelukast in the primary outcome measure FEV<sub>1</sub>, but a small reduction in as-needed short-acting

beta<sub>2</sub>-agonist use (.33 puffs/day) in favor of LTRA was found. No difference was found for worsening asthma, asthma attacks, or quality of life (Simons et al. 2001). Studies of the addition of theophylline to inhaled corticosteroids in children 6 years of age showed both a benefit in quality of life and no benefit (Meltzer et al. 2001). Neither of these theophylline studies had enough quality to generate a recommendation. There is only one study on adjunctive therapy that includes children as young as 4 years of age, and the only studies in children younger than 4 years of age.

#### Step 4: Severe Persistent Asthma

Patients with severe persistent asthma may require high doses of inhaled corticosteroids, a long-acting inhaled beta<sub>2</sub>-agonist and, if needed, an oral corticosteroid (Evidence B). It is the opinion of the Expert Panel that consultation with an asthma specialist is recommended for patients with severe persistent asthma. Evidence to date does not support using a third long-term-control medication added to inhaled corticosteroids and long-acting inhaled beta<sub>2</sub>-agonists in order to avoid using systemic corticosteroid therapy (Evidence C). A study found no benefit for the addition of an LTRA to high doses of inhaled corticosteroids and, for most patients in the study, another medication (either theophylline, a long-acting beta<sub>2</sub>-agonist, oral corticosteroid, or a combination) (Robinson et al. 2001). Similar studies of other long-term-control medications added to the combination of medium-to-high doses of inhaled corticosteroids and long-acting inhaled beta<sub>2</sub>-agonists in severe persistent asthma are not available.

Patients whose asthma is not controlled on high doses of inhaled corticosteroids and the addition of long-acting inhaled beta<sub>2</sub>-agonists also will need oral systemic corticosteroids on a regularly scheduled, long-term basis. For patients who require long-term systemic corticosteroids:

- Use the lowest possible dose (single dose daily or, preferably, on alternate days).
- Monitor patients closely for corticosteroid adverse side effects (see component 3-Medications).

- When control of asthma is achieved, make persistent attempts to reduce systemic corticosteroids. High doses of inhaled corticosteroids are preferable to systemic corticosteroids because inhaled corticosteroids have fewer systemic effects.
- Recommend consultation with an asthma specialist.

#### Recommendations for Future Research

The Panel recommends the following research to clarify treatment options:

Long-term studies to examine the effect of adjunctive therapy on possible loss in pulmonary function and the natural history of asthma—hospitalization, exacerbations, and decline in pulmonary function.

Studies of noninvasive markers that would give a composite picture of both disease activity (e.g., inflammation) and disease control. These could be used as surrogate markers for overall asthma control to guide therapy. Ideally, such markers would be more efficient than gauging a patient's response to therapy following a relatively long therapeutic trial.

Long-term studies to examine the importance of the greater suppression of inflammation achievable with higher doses of inhaled corticosteroids compared with adjunctive therapy. Low doses of inhaled corticosteroids usually are sufficient for improvement in lung function and control of asthma symptoms but may not suppress inflammation to the same extent as higher doses. Studies to assess the value of maximum suppression of inflammation vis-à-vis therapeutic control will contribute to understanding the appropriate use of inhaled corticosteroids and adjunctive therapy.

- Evaluations of adjunctive therapies in children younger than 12 years of age.

## Key Evidence Tables

**Table 1-6. Meta-Analysis: Lung Function Outcomes for Studies Comparing the Addition of Long-Acting Beta<sub>2</sub>-Agonists to a Fixed Dose of Inhaled Corticosteroids**

Meta-Analysis	Effect Size Estimate	95% CI	Test for Homogeneity P-Value	Treatment Effect Estimate	95% CI
FEV <sub>1</sub> : Combined Studies (n = 14)	0.334	0.241, 0.428	0.10	0.17 L 3.71% pred	0.12, 0.22 2.67, 4.75
FEV <sub>1</sub> : Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment and meet most (>4) asthma-specific criteria (n = 3)	0.319	0.139, 0.499	0.14	0.17 L 3.43% pred	0.07, 0.26 1.54, 5.54
FEV <sub>1</sub> : Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment (N = 11)	0.368	0.257, 0.478	0.20	0.19 L 4.08% pred	0.13, 0.25 2.85, 5.30
PEF: Combined studies (n = 9)	0.581	0.417, 0.745	0.0034	24.68 L/min 7.26% pred	17.70, 31.65 5.21, 9.31
PEF: Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment and meet most (>4) asthma-specific criteria (n = 4)	0.643	0.460, 0.826	0.17	27.33 L/min 8.04% pred	19.55, 35.10 5.75, 10.32
PEF: Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment (n = 8)	0.630	0.478, 0.781	0.06	26.77 L/min 7.88% pred	20.32, 33.19 5.98, 9.76

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-E044. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

**Table 1-7. Meta-Analysis: Medication Use Outcomes for Studies Comparing the Addition of Long-Acting Beta<sub>2</sub>-Agonists to a Fixed Dose of Inhaled Corticosteroids**

Meta-Analysis	Treatment Effect Estimate	95% CI	Test for Homogeneity P-Value
Puffs/day: Combined studies (n = 6)	-1.18	-1.56, -0.80	0.018
Puffs/day: Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment and meet most (>4) asthma-specific criteria (n = 3)	-1.34	-1.87, -0.84	0.20
Puffs/day: Sensitivity analysis by quality: Studies meet all generic quality criteria except allocation concealment (n = 5)	-1.00	-1.34, -0.66	0.14

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-E044. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.



Table 1–8. Meta-Analysis: Lung Function Outcomes for Studies Comparing a Lower Dose of Inhaled Corticosteroids Plus Long-Acting Inhaled Beta<sub>2</sub>-Agonist\* vs. Increased Dose of Inhaled Corticosteroids

Meta-Analysis	Effect Size Estimate	95% CI	Test for Homogeneity P-Value	Treatment Effect Estimate	95% CI
FEV <sub>1</sub> : Combined Studies (n = 8)	0.209	0.133, 0.285	0.93	0.11 L 2.32% pred	0.07, 0.15 1.48-3.16
FEV <sub>1</sub> : Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment and meet most (>4) asthma-specific criteria (n = 4)	0.203	0.107, 0.299	0.94	0.11 L 2.25% pred	0.06, 0.16 1.19, 3.32
FEV <sub>1</sub> : Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment (n = 7)	0.212	0.134, 0.290	0.88	0.11 L 2.35% pred	0.07, 0.15 1.49, 3.22
PEF: Combined studies (n = 10)	0.310	0.192, 0.429	0.0002	11.6 L/min 3.4% pred	5.2-18.0 1.5-5.3
PEF: Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment and meet most (>4) asthma-specific criteria (n = 4)	0.300	0.030, 0.569	0.000007	12.75 L/min 3.75% pred	1.28, 24.18 0.38, 7.11
PEF: Sensitivity analysis by quality: Studies that meet all generic quality criteria except allocation concealment (n = 7)	0.296	0.143, 0.449	0.00005	12.58 L/min 3.7% pred	6.08, 19.08 1.79, 5.61

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

## References

- Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.
- Busse W, Nelson H, Wolfe J, Kalberg C, Yancey SW, Rickard K. Comparison of inhaled salmeterol and oral zafirlukast in patients with asthma. *J Allergy Clin Immunol* 1999;103(6):1075-80.
- Davies B, Brooks G, Devoy M. The efficacy and safety of salmeterol compared to theophylline: meta-analysis of nine controlled studies. *Respir Med* 1998;92(2):256-63.
- Eickelberg O, Roth M, Lox R, Bruce V, Rudiger J, Johnson M, Block LH. Ligand-independent activation of the glucocorticoid receptor by beta<sub>2</sub>-adrenergic receptor agonists in primary human lung fibroblasts and vascular smooth muscle cells. *J Biol Chem* 1999; Jan 8;274(2):1005-10.
- Evans DJ, Taylor DA, Zetterstrom O, et al. A comparison of low-dose inhaled budesonide plus theophylline and high-dose inhaled budesonide for moderate asthma. *N Engl J Med* 1997;337(20):1412-8.
- Fjellbirkeland L, Gulsvik A, Palmer JB. The efficacy and tolerability of inhaled salmeterol and individually dose-titrated, sustained-release theophylline in patients with reversible airways disease. *Respir Med* 1994;88(8):599-607.
- Greening AP, Ind PW, Northfield M, Shaw G. Added salmeterol versus higher-dose corticosteroid in asthma patients with symptoms on existing inhaled corticosteroid. *Lancet* 1994;344(8917):219-24.
- Lavolette M, Malmstrom K, Lu S, Chervinsky P, Pujet JC, Peszek I, Zhang J, Reiss TF. Montelukast added to inhaled beclomethasone in treatment of asthma. Montelukast/Beclomethasone Additivity Group. *Am J Respir Crit Care Med* 1999;160(6):1862-8.
- Lazarus SC, Boushey HA, Fahy JV, Chinchilli VM, Lemanske Jr RF, Sorkness CA, Kraft M, Fish JE, Peters SP, Craig T et al. Long-acting  $\beta_2$ -agonist monotherapy vs. continued therapy with inhaled corticosteroids in patients with persistent asthma: a randomized controlled trial. *JAMA* 2001;285(20):2583-93.
- Lemanske Jr RF, Sorkness CA, Mauger EA, Lazarus SC, Boushey HA, Fahy JV, Drazen JM, Chinchilli VM, Craig T, Fish JE, et al. Inhaled corticosteroid reduction and elimination in patients with persistent asthma receiving salmeterol: a randomized controlled trial. *JAMA* 2001;285(20):2594-603.
- Lim S, Jatakanon A, Gordon D, Macdonald C, Chung KF, Barnes PJ. Comparison of high dose inhaled steroids, low dose inhaled steroids plus low dose theophylline, and low dose inhaled steroids alone in chronic asthma in general practice. *Thorax* 2000;55(10):837-41.
- Lofdahl CG, Reiss TF, Leff JA, Israel E, Noonan MJ, Finn AF, Seidenberg BC, Capizzi T, Kundu S, Godard P. Randomised, placebo controlled trial of effect of a leukotriene receptor antagonist, montelukast, on tapering inhaled corticosteroids in asthmatic patients. *BMJ* 1999;319(7202):87-90.
- McIvor RA, Pizzichini E, Turner MO, Hussack P, Hargreave FE, Sears MR. Potential masking effects of salmeterol on airway inflammation in asthma. *Am J Respir Crit Care Med* 1998;158(3):924-30.
- Meltzer EO, Orgel HA, Ellis EF, Eigen HN, Hemstreet MP. Long-term comparison of three combinations of albuterol, theophylline, and beclomethasone in children with chronic asthma. *J Allergy Clin Immunol* 1992 Jul;90(1):2-11.
- Muir JF, Bertin L, Georges D. French Multicentre Study Group. Salmeterol versus slow release theophylline combined with ketotifen in nocturnal asthma: a multicenter study. *Eur Respir J* 1992;5(10):1197-200.
- Nassif EG, Weinberger M, Thompson R, Huntley W. The value of maintenance theophylline in steroid-dependent asthma. *N Engl J Med* 1981; Jan 8;304(2):71-5.



- Nelson HS, Busse WW, Kerwin E, Church N, Emmett A, Richard K, Knobil K. Fluticasone propionate/salmeterol combination provides more effective asthma control than low-dose inhaled corticosteroid plus montelukast. *J Allergy Clin Immunol* 2000;106(6):1088-95.
- Paggiaro PL, Giannini D, Di Franco A, Testi R. European Study Group. Comparison of inhaled salmeterol and individually dose-titrated slow-release theophylline in patients with reversible airway obstruction. *Eur Respir J* 1996;9(8):1689-95.
- Pauwels RA, Lofdahl CG, Postma DS, Tattersfield AE, O'Byrne P, Barnes PJ, Ullman A. Formoterol and Corticosteroids Establishing Therapy (FACET) International Study Group. Effect of inhaled formoterol and budesonide on exacerbations of asthma. *N Engl J Med* 1997;337(20):1405-11.
- Pierson WE, LaForce CF, Bell TD, MacCosbe PE, Sykes RS, Tinkelman D. Long-term, double-blind comparison of controlled-release albuterol versus sustained-release theophylline in adolescents and adults with asthma. *J Allergy Clin Immunol* 1990;85(3):618-26.
- Roberts JR, Desai HI, Gillespie CA, Simons FE. Sustained-release terbutaline vs. sustained-release theophylline in young patients with asthma. *Am J Dis Child* 1986;140(7):650-54.
- Russell G, Williams DA, Weller P, Price JF. Salmeterol xinafoate in children on high dose inhaled steroids. *Ann Allergy Asthma Immunol* 1995;75(5):423-8.
- Shrewsbury S, Pyke S, Britton M. Meta-analysis of increased dose of inhaled steroid or addition of salmeterol in symptomatic asthma (MIASMA). *BMJ* 2000;320(7246):1368-73.
- Simons FE, Villa JR, Lee BW, Teper AM, Lyttle B, Aristizabal G, Laessig W, Schuster A, Perez-Frias et al. Montelukast added to budesonide in children with persistent asthma: a randomized, double-blind, crossover study. *J Pediatr* 2001;138(5):694-8.
- Sont JK, Willems LN, Bel EH, van Krieken JH, Vandenbroucke JP, Sterk PJ. The AMPUL Study Group. Clinical control and histopathologic outcome of asthma when using airway hyperresponsiveness as an additional guide to long-term treatment. *Am J Respir Crit Care Med* 1999;159(4 Pt 1):1043-51.
- Ukena D, Harnest U, Sakalauskas R, Magyar P, Vetter N, Steffen H, Leichtl S, Rathgeb F, Keller A, Steinijans VW. Comparison of addition of theophylline to inhaled steroid with doubling of the dose of inhaled steroid in asthma. *Eur Respir J* 1997;10(12):2754-60.
- Verberne AA, Frost C, Duiverman EJ, Grol MH, Kerrebijn KE. Addition of salmeterol versus doubling the dose of beclomethasone in children with asthma. The Dutch Asthma Study Group. *Am J Respir Crit Care Med* 1998;158(1):213-9.
- Weinstein SF, Pearlman DS, Bronsky EA, Byrne A, Arledge T, Liddle R, Stahl E. Efficacy of salmeterol xinafoate powder in children with chronic persistent asthma. *Ann Allergy Asthma Immunol* 1998;81(1):51-8.
- Woolcock A, Lundback B, Ringdal N, Jacques LA. Comparison of addition of salmeterol to inhaled steroids with doubling of the dose of inhaled steroids. *Am J Respir Crit Care Med* 1996;153(5):1481-8.

## Use of Antibiotics To Treat Asthma Exacerbations

### Question

***Does routinely adding antibiotics to standard care improve the outcomes of treatment for acute exacerbation of asthma? Does the addition of antibiotics to standard care in the following populations improve the outcomes of treatment for an acute exacerbation of asthma: patients without signs and symptoms of bacterial infection; patients with signs and symptoms of a bacterial infection; patients with signs and symptoms of sinusitis?***

### Summary Answer to the Question

The available evidence (two randomized, controlled clinical trials) suggests no benefit from antibiotic therapy for asthma exacerbations, whether administered routinely or when suspicion of bacterial infection is low (SRE-Evidence B). No studies addressed the question of greatest relevance to contemporary clinical practice: whether the addition of antibiotics to standard care when signs and symptoms suggest the possibility—but do not clearly indicate the presence—of bacterial infection improves the outcomes of treatment for acute asthma exacerbations.

The EPR-2 recommendation has not been changed: Antibiotics are not recommended for the treatment of acute asthma exacerbations except as needed for comorbid conditions—e.g., for the patients with fever and purulent sputum, evidence of pneumonia, or suspected bacterial sinusitis.

### Rationale for the Question

Asthma exacerbations often are associated with clinical signs of infection, such as purulence of expectorated sputum or nasal discharge. Most asthma exacerbations are associated with infection by a respiratory virus, especially rhinovirus (Nicholson et al. 1993; Johnston et al. 1995), but a small percentage of exacerbations are associated with

infection by an atypical bacterium, like *Mycoplasma pneumoniae* or *Chlamydia pneumoniae* (Freymuth et al. 1999). It is widely believed that coincident bacterial sinusitis contributes to asthma exacerbations, and some clinicians have postulated that airway obstruction due to mucus plugging—common in asthma—predisposes patients to bacterial infection of nondraining regions of the lungs.

In the absence of clear signs of bacterial infection (e.g., lobar pulmonary infiltrate on chest radiography distinguishing viral from bacterial infections), infection is often difficult to manage. Viral infections commonly resemble bacterial infections in that they also cause neutrophilic inflammation of the upper and lower airways (Teran et al. 1997; Trigg et al. 1996; Fahy et al. 1995). This difficulty, coupled even with the remote possibility that bacterial infection may be associated with an asthma exacerbation, may account for the frequency with which antibiotics are prescribed in addition to inhaled bronchodilators, inhaled or systemic corticosteroids, and supplemental oxygen.

### Summary of the Evidence

The following description of the SRE is an adaptation of the evidence report, including direct excerpts, submitted by the Blue Cross Blue Shield Association Evidence-Based Practice Center. (See Introduction, Methods.)

### Methods of Literature Search

In addition to the selection criteria for studies related to all topics in the SRE (described in the Introduction section), studies for this question were included in which standard care (asthma medications) plus antibiotics was compared with standard care alone in the treatment of acute asthma exacerbations. Patient populations included patients without signs and symptoms of bacterial infection, patients with signs and symptoms of bacterial infection, and patients with signs and symptoms of sinusitis.

## I Summary of Findings

### Studies

Only two randomized, double-blind, placebo-controlled, parallel-group trials—with a total enrollment of 121 patients—have addressed the question of whether routinely adding antibiotics to standard care improves the outcomes of treatment for acute asthma exacerbations (Shapiro et al. 1974; Graham et al. 1982). (See the key evidence tables in this section.) Both trials studied patients hospitalized for asthma exacerbations. Both used a penicillin derivative whose activity against atypical bacteria was unknown. Shapiro and colleagues examined the effects of hetacillin (an analogue of ampicillin; 100 mg/kg every 24 hours for a minimum of 24 hours, then 225 mg four times per day for 6 days) in 50 children who did not exhibit clinical evidence of bacterial infection. Graham and colleagues examined the effects of amoxicillin (500 mg three times per day) in 60 adults and adolescents who experienced a total of 71 hospital admissions. Whereas the pediatric study explicitly excluded patients with clinical evidence of bacterial infection, the study of adults and adolescents excluded only patients with evidence of pneumonia on chest radiography. Thus, the populations in these studies consisted primarily of patients without signs or symptoms of bacterial illness, including suspected acute sinusitis.

In both trials, all patients received standard care that included high-dose oral or intravenous corticosteroids and regularly scheduled beta<sub>2</sub>-agonist treatment. In the pediatric study, all patients were also treated with intravenous aminophylline followed by oral theophylline.

The study design and conduct for these two trials did not meet the SRE criteria for higher quality because of deficiencies in allocation concealment, subject withdrawal, and reporting of power calculations.

The outcomes analyzed included change in FEV<sub>1</sub>, symptom scores, and length of hospital stay.

### I Results of Studies

Neither study reported an association—nor a trend towards an association—between antibiotic treatment

and greater improvement in any asthma outcome. Therefore, available evidence suggests no benefit from the use of antibiotic treatment for asthma exacerbations either routinely or when the suspicion of bacterial infection is minimal. (See key evidence tables 1–11 and 1–12.)

### Additional Literature/Information

A related question, for which clinical trials data are unavailable, should ask whether the use of an antibiotic active against *Mycoplasma* and *Chlamydia* would alter outcomes. Some recent studies using polymerase chain reaction (PCR)-based methods for detecting specific genomic sequences have suggested that chronic infection with these organisms may contribute to the severity of chronic asthma (Kraft et al. 1998). These highly sensitive methods have not yet been applied to the analysis of airway tissue or secretions obtained from patients suffering acute exacerbations. Thus, there is a theoretical basis for the concept that a subgroup of patients with asthma exacerbations may benefit from treatment with an antibiotic that is active against these atypical bacteria.

The EPR-2 statement that “the use of antibiotics is generally reserved for patients with fever and purulent sputum (discolored because of polymorphonuclear leukocytes, not eosinophils)” comes under scrutiny because low-grade fever also may accompany viral respiratory infections. Furthermore, a recent study shows that discoloration of sputum by polymorphonuclear leukocytes is observed in viral tracheobronchitis, and the sputum from patients suffering from uncomplicated asthma exacerbations commonly contains high numbers of polymorphonuclear leukocytes (Fahy et al. 1995).

### Recommendations for EPR Update

No evidence supports changing the EPR-2 recommendation (SRE-Evidence B). The parenthetical statement on page 116 of EPR-2 [“(discolored because of polymorphonuclear leukocytes, not eosinophils)”] should be removed (Evidence C). The recommendation can otherwise stand and is as follows:

Antibiotics are *not* recommended for the treatment of acute asthma exacerbations except as needed for comorbid conditions. Bacterial, *Chlamydia*, or *Mycoplasma* infections infrequently contribute to exacerbations of asthma and therefore the use of antibiotics is generally reserved for patients with fever and purulent sputum and for patients with evidence of pneumonia. When the presence of bacterial sinusitis is suspected, treat with antibiotics.

#### Recommendations for Future Research

No studies addressed the question of greatest relevance to contemporary clinical practice—whether the addition of antibiotics to standard care when signs and symptoms suggest the possibility but do not clearly indicate the presence of bacterial infection improves the outcomes of treatment for acute asthma exacerbations. The two trials reviewed excluded the patients most likely to be treated with antibiotics and those with signs or symptoms suggestive of bacterial infection, including suspected acute sinusitis. Studies of the efficacy of antibiotic treatment in this group are needed.

Several studies are needed to clarify the role of antibiotics in the treatment of asthma exacerbations. Questions for research are as follows:

- What is the efficacy of antibiotic treatment in asthma patients most likely to be treated with antibiotics, such as those with signs suggestive of bacterial infection, including suspected acute sinusitis? The role of sinusitis in acute exacerbations of asthma has not been truly defined.
  - What is the role of sinusitis in acute exacerbations of asthma or increased asthma severity?
  - What is the efficacy of using an antibiotic active against atypical bacteria, given the possibility that such bacteria commonly contribute to asthma exacerbations?
  - What would be the value of studies applying modern sensitive methods of detection of atypical bacteria (e.g., PCR-based methods) to samples of airway tissues or secretions obtained at the time of an asthma exacerbation?
- Do antibiotics such as macrolides have a non-antibiotic action (e.g., anti-inflammatory) that is beneficial in asthma patients?

## Key Evidence Tables

Table 1-9. Study Characteristics

Citation	Study Design	Study Setting	Asthma Severity	Eligibility
Graham, Milton, Knowles et al. 1982	Randomized, double-blind, placebo-controlled, parallel group trial	Country: United Kingdom  Funding: Government grant  Tx setting: University Hospital, inpatient setting	Stated: Not specified  Estimated: Unable to estimate	Eligibility assessed on admission to hospital with asthma exacerbation:  <ul style="list-style-type: none"> <li>• FEV<sub>1</sub> of 1.5L or less and/or PEF of 150 l/min</li> <li>• Reversibility of FEV<sub>1</sub> at least 15% spontaneously or after inhalation of beta<sub>2</sub>-agonist</li> </ul> Exclusions: Evidence of pneumonia on CXR, history of penicillin allergy
Shapiro, Eggleston, Pierson et al. 1974	Randomized, double-blind, placebo-controlled, parallel group trial	Country: United States  Funding: Pharm Industry and Government grant  Tx setting: Hospital, inpatient setting	Stated: Not specified  Estimated: Unable to estimate	Eligibility assessed on admission to hospital with asthma exacerbation:  <ul style="list-style-type: none"> <li>• Severe bronchospasm, lack of response to subcutaneous epinephrine</li> </ul> Exclusions: Clinical evidence of bacterial infection; recent use of antibiotics

## Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. *Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44*. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.



Table 1–10. Study Parameters

Citation	Study Arm	Treatment	Comments
Graham, Milton, Knowles et al. 1982	Placebo	Placebo tablet 3 times per day  Oral prednisolone (20–60 mg/day) and/or IV hydrocortisone (100–200 mg every 4 to 6 hours)  Regularly scheduled beta <sub>2</sub> -agonists and/or phosphodiesterase inhibitors  Chest physiotherapy	60 patients enrolled with 71 exacerbations. Unit of analysis by exacerbations.          Culture-proven bacterial source of infection found in two patients on admission and two patients on discharge
	Antibiotics	Amoxicillin 500 mg 3 times per day  Oral prednisolone (20–60 mg/day) and/or IV hydrocortisone (100–200 mg every 4 to 6 hours)  Regularly scheduled beta <sub>2</sub> -agonists and/or phosphodiesterase inhibitors  Chest physiotherapy	
Shapiro, Eggleston, Pierson et al. 1974	Placebo	Placebo 4 times per day for 6 days  IV hydrocortisone (7 mg/kg/24 hr) for 24 hours, followed by oral prednisone  IV aminophylline (15 mg/kg/24 hr) for 24 hours, followed by oral theophylline  Nebulized beta <sub>2</sub> -agonists q30 min x 4, then as needed	37 patients enrolled with 44 exacerbations, unit of analysis by exacerbation
	Antibiotics	Hetacillin (100 mg/kg/24 hr) for at least 24 hours, followed by oral hetacillin 225 mg 4 times per day for 6 days  IV hydrocortisone (7 mg/kg/24 hr) for 24 hours, followed by oral prednisone  IV aminophylline (15 mg/kg/24 hr) for 24 hours, followed by oral theophylline  Nebulized beta <sub>2</sub> -agonists q30 min x 4, then as needed	

**Source:**

Blue Cross and Blue Shield Association Technology Evaluation Center. *Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44*. AHRQ Publication No. 01–EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

## Key Evidence Tables

Table 1-11. Lung Function Outcome

Citation	Study Arm	Number Enrolled	Number Evaluable	Study Duration (days) (median/range)	FEV <sub>1</sub> Baseline (mean)
Graham, Milton, Knowles et al. 1982	Placebo	71*	32*	8 (3-16)	20.9 (<7.3-63)
	Antibiotics	71*	37*	7 (3-25)	23.1 (<7.3-45.5)
Shapiro, Eggleston, Pierson et al. 1974	Placebo	50*	24*	2.9 (SD 1.4)	26.5 (SD 15)
	Antibiotics	50*	20*	2.5 (SD 0.8)	28.3 (SD 11)

Table 1-12. Symptoms/Utilization Outcome

Citation	Study Arm	Number Enrolled	Number Evaluable	Study Duration (days) (median/range)
Graham, Milton, Knowles et al. 1982	Placebo	34	32	8 (3-16)
	Antibiotics	37	37	7 (3-25)
Shapiro, Eggleston, Pierson et al. 1974	Placebo	24	24	2.9 (SD 1.4)
	Antibiotics	20	20	2.5 (SD 0.8)

	FEV <sub>1</sub> Final (mean)	P-Value	PEF Baseline (mean/range)	PEF Final (mean/range)	P-Value
	65.6 (31.5–108.5)		23.8 (<9.4–83.9)	72.8 (32.8–108.1)	
	52.3 (10–92.9)	0.039	23.8 (<9.4–47.3)	59 (16.7–95)	0.052
	49 (SD 17)		NR	NR	
	61 (SD 19)	NR	NR	NR	

\*Unit of analysis was admission. Number enrolled represented total admissions in both groups, information not provided by group. Number evaluated represents total number of admissions included in analysis.

**Source:**

Blue Cross and Blue Shield Association Technology Evaluation Center. *Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44*. AHRQ Publication No. 01–EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

	Baseline Symptom Score (median/range)	Final Symptom Score (median/range)	P-Value	Hospital Length of Stay	P-Value
	11 (6–12)	4 (4–8)		8 (3–16)	
	11 (5–12)	5 (4–9)	NS	7 (3–25)	NS
	7.1 (mean) (SD 2.2)	2.5 (SD 2.0)		2.9 (SD 1.4)	
	7.1 (mean) (SD 1.8)	2.0 (SD 2.0)	NR	2.4 (SD 0.8)	

**Source:**

Blue Cross and Blue Shield Association Technology Evaluation Center. *Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44*. AHRQ Publication No. 01–EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

## References

- Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.
- Fahy JV, Jim KW, Liu J, Boushey HA. Prominent neutrophilic inflammation in sputum from subjects with asthma exacerbation. *J Allergy Clin Immunol* 1995;95(4):843-52.
- Freyndt F, Vabret A, Brouard J, Toutain F, Verdon R, Petjean J, Gouarin S, Duhamel JF, Guillois B. Detection of viral, Chlamydia pneumoniae and Mycoplasma pneumoniae infections in exacerbations of asthma in children. *J Clin Virol* 1999;13(3):131-9.
- Graham VA, Milton AF, Knowles GK, Davies RJ. Routine antibiotics in hospital management of acute asthma. *Lancet* 1982;1(8269):418-420.
- Johnston SL, Pattermore PK, Sanderson G, Smith S, Lampe F, Josephs L, Symington P, O'Toole S, Myint SH, Tyrell DA, et al. Community study of role of viral infections in exacerbations of asthma in 9-11 year old children. *BMJ* 1995;310(6989):1225-9.
- Kraft M, Cassell GH, Henson JE, Watson H, Williamson J, Marmion BP, Gaydos CA, Martin RJ. Detection of Mycoplasma pneumoniae in the airways of adults with chronic asthma. *Am J Respir Crit Care Med* 1998;58(3):998-1001.
- Labro MT. Anti-inflammatory activity of macrolides: a new therapeutic potential? *J Antimicrob Chemother* 1998;41(Suppl B):37-46.
- Nicholson K, Kent J, Ireland D. Respiratory viruses and exacerbations of asthma in adults. *Br Med J* 1993;307:982-6.
- Shapiro GG, Eggleston PA, Pierson WE, Ray CG, Bierman CW. Double-blind study of the effectiveness of a broad spectrum antibiotic in status asthmaticus. *Pediatrics* 1974;53(6):867-72.
- Teran LM, Johnston SL, Schroder JM, Church MK, Holgate ST. Role of nasal interleukin-8 in neutrophil recruitment and activation in children with virus-induced asthma. *Am J Respir Crit Care Med* 1997;155(4):1362-6.
- Trigg CJ, Nicholson KG, Wang JH, Ireland DC, Jordan S, Duddle JM, Hamilton S, Davies RJ. Bronchial inflammation and the common cold: a comparison of atopic and non-atopic individuals. *Clin Exp Allergy* 1996;26(6):665-76.



Expert Panel Report  
Guidelines for the  
Diagnosis and  
Management of  
Asthma

## 2 Monitoring

Asthma  
Reference



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## 2. Monitoring

Two distinct questions have been raised regarding the use of written action plans in the management of asthma. First, does the use of written action plans make a difference in patient outcomes beyond those accomplished by appropriate medical/pharmacologic management? Second, is there a difference in patient outcomes between action plans based on symptom monitoring and those based on peak flow monitoring? This section of the EPR Update considers both questions.

### Written Action Plans Compared to Medical Management Alone

#### Question

***Compared to medical management alone, does the use of a written asthma action plan improve outcomes?***

#### Summary Answer to the Question

Data are insufficient to support or refute the benefits of using written asthma action plans compared to medical management alone (SRE-Evidence B). Seven studies compared medical management with written action plans to medical management without action plans. Beyond including instructions on the action plan to the intervention groups, four of these studies did not include asthma education for either the intervention or control groups; three of the studies included similar but limited asthma education for both intervention and control groups. Only one study included children. Significant limitations in study designs and methods in these studies preclude conclusions. For example, the studies showing no benefits of written action plans did not have sufficient power for comparisons between treatment and control groups, and the two studies reporting significant improvements with action plans had potential biases in patient

selection, withdrawals, data collection, or analysis.

However, a Cochrane review of 25 studies comparing asthma self-management education interventions for adults to medical care without such education also contrasted those studies with self-management interventions that included written action plans to those that did not. The self-management interventions that included written action plans had the greatest benefits, including reduced emergency department visits and hospitalizations and improved lung function.

The EPR-2 recommendations have not been changed: It is the opinion of the Expert Panel that use of written action plans as part of an overall effort to educate patients in self-management is recommended, especially for patients with moderate or severe persistent asthma and patients with a history of severe exacerbations (Evidence B, C).

#### Other Question

The use of written action plans is recommended in the EPR-2 and is widely accepted as good practice. Generally, the use of written action plans has been studied as part of self-management education (Gibson et al. 2000). In busy practices, however, physicians often provide their patients with action plans independent of other asthma education efforts. This question was posed in order to identify data that describe the effects of using written action plans, independent of other components of asthma education.



## Systematic Review of the Evidence

The following description of the SRE is an adaptation of the evidence report, including direct excerpts, submitted by the Blue Cross Blue Shield Association Evidence-Based Practice Center. (See Introduction, Methods.)

### I Methods of Literature Search

For the purpose of the SRE, an action plan is a written algorithm that identifies specific clinical indicators that should alert patients to make adjustments in their medications and provides specific instructions on how to make these adjustments. EPR-2 recommends the use of both a daily self-management plan and an action plan for exacerbations. Generally, studies included in the SRE involved the use of one plan that combined the objectives of both. Typically, the plans divided steps for patient actions into different zones, in which recommended actions are correlated with differing acute signs and symptoms of worsening asthma. Most of the plans in the available studies used four-zone plans, some were three-zone plans that did not include directions for use of oral corticosteroids before seeking emergency care.

The evidence review examined studies in which the intervention used an action plan as defined above and, if asthma education was given to both treatment and control groups, the treatment group had no more than 1 additional hour of education for the action plan. The treatment/observation duration was at least 12 weeks, and the intervention and control groups received the same treatment, except that the intervention group also received a written action plan. Studies were excluded if the comparisons were confounded by additional treatment components in the intervention group—for example, optimization of medications in the intervention group only or education programs of more than 1 hour in the intervention group only. The literature review included randomized controlled trials (RCTs) in which at least 25 evaluable patients (not physicians) were randomly allocated to the intervention and control groups.

## I Summary of Findings

### Studies

Seven studies involving more than 1,400 patients met SRE inclusion criteria for review; only one of the studies included children. (See the key evidence tables in this section.) None of the studies met SRE standards for high quality; each had significant limitations. None was conducted with sufficient power (i.e., adequate numbers of subjects in each study arm) to enable comparisons between treatment and control groups. In one study reporting reduced emergency department visits, data were unavailable to control for baseline differences that may have existed between treatment and control groups, and the reported effect may be attributed to a subset of high frequency users. In another study, the design involved clinicians who both provided plans and collected assessment data. Moreover, a large number of subjects were excluded from the analyses.

All seven studies compared medical management with written action plans to medical management without written action plans, and all used a peak flow meter-based plan. Three of the studies also included similar but limited asthma education for both the intervention and control groups, but the groups still differed as to whether written plans were used. In two trials, the control group used peak flow meters but without an action plan.

### Results of Studies

Five trials documented no differences in outcomes, and two trials documented significant benefit of written action plans, especially in reducing emergency department visits. However, there were notable limitations to each of these trials, as described earlier. In summary, SRE study data were insufficient to support or to refute the advantages of using asthma action plans independent of self-management education when compared with medical management alone.

## Additional Literature/Information

Evidence supporting the use of written plans as a component of self-management education is reported in a recent Cochrane Collaboration review (Gibson et al. 2000). The SRE question on action plans provides a clearer assessment of isolating the advantages of providing an action plan. The Cochrane review centered on the benefits of self-management interventions and regular medical review with the clinician vs. usual medical care. The Cochrane review, however, also contrasted those self-management interventions with written action plans to those without written action plans. The review included some of the same studies included in the SRE but overcame the limitations of study sample sizes by pooling data. Further, the set of 25 studies in the Cochrane review was larger than the 7 in the SRE due to the broader question under review. In the Cochrane analysis that compared results of self-management interventions with action plans to those without, the interventions with written action plans demonstrated the greatest benefits, including reduced asthma-related hospital admissions (odds ratio 0.35, 95 percent confidence interval) and reduced emergency department visits (odds ratio 0.55, 95 percent confidence interval). In addition, patients who managed their asthma by adjusting medications according to a written action plan had better lung function than those whose medications were adjusted by a doctor during regular care visits. The review concluded that training in asthma self-management that involves self-monitoring by either peak flow or symptoms, coupled with regular medical review and a written action plan, appears to improve health outcomes for adults with asthma.

Additional evidence supporting written action plans coupled with regular patient education and medical review is available from a recent case control study (Abramson et al. 2001). This study does not fit the SRE review criteria because studies that qualified for this review were required to be RCTs allowing inferences of cause and effect, and they were required to provide an action plan independent of a multicomponent intervention including education. Although the Abramson study is not an RCT, it is a well-conducted study that compared 51 patients who died from asthma to 202 patients presenting to hospitals with acute

asthma. The study reported that written action plans for patients with severe persistent asthma were associated with a 70 percent reduction in mortality risk. As such, the study supports the opinion that providing written action plans as part of asthma education is an important element of practice.

## Recommendations for EPR Update

No data from the SRE, in which RCTs compared written action plans to medical management alone, indicate the need to change the EPR-2 action plan recommendations (SRE-Evidence B). Additional data from studies on action plans as a part of self-management education support the EPR-2 recommendations (Evidence B, C). <sup>1</sup> Following blue text indicates revisions that should be incorporated into the text on pages 33 and 123 in EPR-2.

Component 1: Measures of Assessment and Monitoring; Periodic Assessment and Monitoring (page 33 in EPR-2)

Whether peak flow monitoring, symptom monitoring, or a combination of approaches is used, the Expert Panel believes that self-monitoring is important to the effective self-management of asthma. The nature and intensity of self-monitoring should be individualized, based on such factors as asthma severity, patient's ability to perceive airflow obstruction, availability of peak flow meters, and patient preferences.

It is the opinion of the Expert Panel that, regardless of the type of monitoring used, patients should be given a written action plan and instructed to use it. (See figure 4–5.) It is the opinion of the Expert Panel that including action plans as part of an overall effort to educate patients in self-management is the soundest approach and is especially indicated for patients with moderate or severe persistent disease or a history of severe exacerbations (Evidence B, C). It is the opinion of the Expert Panel that a plan is important in large part because it enhances clinician-patient communication. The plan should define a regimen that meets the medical needs of the patient

and should have a format that facilitates the patient's understanding and ability to take appropriate action to control the disease. Regardless of format, an effective plan should include the following:

- Explicit, patient-specific instructions for environmental and other preventive efforts that may help to prevent or reduce the impact of exacerbations.
- An algorithm of procedures that describes how to use long-term-control medicines, given a set of specific circumstances and conditions, and clear instructions on how to adjust medicines when conditions change.
- Steps the patient should take when medicines are ineffective or if an emergency situation.
- Contacts for securing urgent care, if needed.

As emphasized above, it is the opinion of the Expert Panel that a written action plan is considered a part of ongoing efforts to provide self-management education and support appropriate to the severity of the patient's asthma, the patient's age, and related circumstances (Evidence B, C). The clinician should periodically review the plan, revise it as necessary, and confirm that the patient knows what to do if his or her asthma gets worse.

#### Component 4: Education for a Partnership in Asthma Care, Key Points (page 123 in EPR-2)

- Patient education should begin at the time of diagnosis and be integrated into every step of clinical asthma care.
- It is essential that education be provided by all members of the health care team. The principal clinician should introduce the key educational messages and negotiate agreements with patients; these messages should be reinforced and expanded by all members of the health care team.
- Teach asthma self-management, tailoring the approach to the needs of each patient. Maintain a sensitivity to cultural beliefs and practices.

- Teach and reinforce at every opportunity:
  - Basic facts about asthma
  - Roles of medications
  - Skills: inhaler/spacer/holding chamber use, self-monitoring
  - Environmental control measures
  - When and how to take rescue actions.
- Jointly develop treatment goals.
- To encourage an active partnership, provide all patients with a written daily self-management plan and an action plan for exacerbations. A written action plan is considered part of ongoing efforts to provide self-management education and support appropriate to the severity of the asthma, the patient's age, and related circumstances (Evidence B, C). Action plans are especially important for patients with moderate-to-severe asthma and patients with a history of severe exacerbations. Provide appropriate patients with a daily asthma diary.
- Encourage adherence by promoting open communication; individualizing, reviewing, and adjusting plans as needed; emphasizing goals and outcomes; and encouraging family involvement.

#### Recommendations for Future Research

Research that may enhance the quality and effect of interventions fostering patient self-management would examine the following questions:

- Are some action plan formats more effective than others? What characterizes the most effective format?
- What alternative action plan formats are effective, given specific patient needs, including disease severity, literacy levels, languages spoken, ages, and unique management problems (e.g., comorbidities)?
- How much time and emphasis should be given to the development of action plans during the course of clinical counseling? In comprehensive education programs? In medical review?



- What are potential means of providing self-management interventions that include action planning to patients who are members of underserved populations (e.g., reaching them through worksites, community centers, or churches)?
- How effective are written action plans in treating children with asthma?
- How effective are written action plans in different caretaker situations (e.g., daycare, camps, or school)?

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## Key Evidence Tables

Table 2-1. Study Characteristics

Citation	Design	Study Setting	
Optimal medical management vs. optimal medical management + peak flow meter (PFM)-based action plan			
Jones, Mullee, Middleton et al. 1995	Randomized; parallel, controlled	Country: United Kingdom Funding: Pharm. ind. grant Tx Setting: Primary/specialty combination, university Multicenter	
Drummond, Abdalla, Beattie et al. 1994 (GRASSIC)	Randomized; parallel, controlled	Country: United Kingdom Funding: Academic grant Tx Setting: Specialty care, nonuniversity Multicenter	
Ayres, Campbell, Follows 1995	Randomized; parallel, controlled	Country: United Kingdom Funding: Pharm. ind. grant Tx Setting: Unknown Multicenter	
Cowie, Revitt, Underwood et al. 1997	Randomized; parallel, controlled	Country: Canada Funding: Hospital Tx Setting: Primary/specialty combination, university Multicenter	
Cote, Cartier, Robichaud et al. 1997	Randomized; parallel, controlled	Country: Canada Funding: Pharm. ind. grant Tx Setting: Specialty care, nonuniversity Multicenter	
Optimal medical management + (PFM) use (without action plan) vs. optimal medical management + PFM-based action plan			
Ignacio-Garcia and Gonzalez-Santos 1995	Randomized; parallel, controlled	Country: Spain Funding: Not specified Tx Setting: Specialty care, nonuniversity	
Charlton, Antoniou, Atkinson et al. 1994	Randomized; parallel, controlled	Country: Australia Funding: Pharm. ind. and government and university funding Tx Setting: Specialty care, nonuniversity	

Eligibility	Comments
<p>Patient eligibility based on symptoms only</p> <p>Included patients using inhaled corticosteroids &lt;1,000 mcg per day for at least 1 month</p> <p>Exclusions: Patients on oral steroids or using peak flow meters at home</p>	<p>Power based on several outcomes (FEV needed 23 patients, sixfold reduction in night wakening needed 21 per group, eightfold reduction in days off work or school needed 37 per group).</p> <p>2-week course of oral steroids given before randomization to optimize lung function.</p>
<p>Patient eligibility based on lung function and utilization</p> <p>Inclusion: FEV<sub>1</sub> reversibility 20% or greater</p> <p>Exclusion: Patients who already owned a PFM</p>	<p>Power based on the 569 randomized, but n varies for each outcome and in some cases is not specified as to exact n, just that n was &gt; = 250; may not be powered for all outcomes.</p> <p>Patients included had less severe asthma on entry than those who already owned a PFM and were excluded, especially with regard to social and physical functioning.</p>
<p>Patient eligibility based on lung function, symptoms, utilization</p> <p>Inclusions: PEF variability maximum 0.15%; nights/week with symptoms minimum 3; use of inhaled corticosteroids or sodium cromoglycate for a minimum of 3 months</p>	<p>Doctor also graded the overall and individual severity of symptoms as 0 = none and 3 = severe.</p>
<p>Patient eligibility based on symptoms and utilization</p> <p>Inclusions: Treatment for an exacerbation of asthma in an ER or attending a university asthma clinic; history of receiving urgent treatment for asthma in the previous 12 months</p>	<p>Subjects were recruited by contacting those who had been treated for an exacerbation of asthma in an emergency room or those attending a university asthma clinic who had a history of having received urgent treatment for their asthma in the previous 12 months.</p>
<p>Patient eligibility based on lung function and symptoms</p> <p>FEV<sub>1</sub> postbronchodilator 85–100% of predicted</p> <p>PEF minimum 85% of predicted; PEF variability minimum 0%;</p> <p>Methacholine</p> <p>Exclusion: Patients having previously taken part in an asthma educational program</p>	<p>In discussion "although the control group received more than the usual care treatment, none received book, none had written action plan; none had structured education or PFM at home after run-in."</p> <p>Run-in = 2–6 wks.; diagnosis of asthma included need to take daily anti-inflammatory agents; were excluded.</p>
<p>Patient eligibility based on utilization only</p> <p>Inclusion: Patients from outpatient asthma clinic with asthma for 2 years</p>	<p>One doctor aware of the group assignment was responsible for assessment of all patients' condition, but the paper also says "in control group, the doctor assessing the patient was blinded with regard to registers of peak flow monitoring until end of study", random allocation by order of recruitment.</p>
<p>Patient eligibility based on utilization only</p> <p>Inclusion: Patients who required admission for asthma or attended the outpatient department</p>	<p>Randomization was based on age, sex, whether they used asthma prophylaxis before study.</p>

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01–EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

Table 2-2. Lung Function Outcomes: FEV<sub>1</sub>

Citation	Study Arm	Number Enrolled	Number Evaluable	Treatment Duration (weeks)	
Usual care vs. peak flow meter-based action plan					
Jones, Mullee, Middleton et al. 1995	Usual care	64	39	26	
	PFM-based action plan	63	33	26	
Drummond, Abdalla, Beattie et al. 1994 (GRASSIC)	Usual care	284	260	52	
	PFM-based action plan	285	250	52	
Ayres, Campbell, Follows 1995	Usual care	64	64	24	
	PFM-based action plan	61	31	24	
Cowie, Revitt, Underwood et al. 1997	Usual care	48			
	PFM-based action plan	46			
Cote, Cartier, Robichaud et al. 1997	Usual care	54			
	PFM-based action plan	50			
Usual care + PFM use alone vs. usual care + PFM-based action plan					
Ignacio-Garcia and Gonzalez-Santo 1995	Usual care + PFM use	44	35	28	
	Usual care + PFM-based action plan	50	35	28	
Charlton, Antoniou, Atkinson et al. 1994	Usual care + PFM use	43			
	Usual care + PFM-based action plan	48			

	Baseline FEV <sub>1</sub> *	FEV <sub>1</sub>	P-Value	P-Value Comparison	Comments
	85.4 +/- 17.5 % of predicted	81.2 +/- 18.3 % of predicted			
	87.1 +/- 16.9 % of predicted	83.2 +/- 18 % of predicted	NS	Absolute value, Tx vs. Ctl	
	78.1 % of predicted	75.4 +/- 27.7 % of predicted			95% CI for baseline FEV is 74.8-81.4.
	77.3 % of predicted	74.6 +/- 27.8 % of predicted	NS	Change, Tx vs. Ctl	95% CI for baseline FEV is 74.1-80.5.
	2 +/- 0.1 L (type predose)	2.2 +/- 0.1 L (type predose)			Unclear number of patients analyzed on each end point.
	2.3 +/- 0.1 L (type predose)	2.3 +/- 0.2 L (type predose)	NS	Absolute value, Tx vs. Ctl	Unclear number of patients analyzed on each end point.
	78 +/- 21.3 % of predicted				Number of subjects with <60% predicted was 10.
	82 +/- 20.5 % of predicted				Number of subjects with <60% predicted was 9.
	65.34 +/- 16.6 % of predicted (type predose)	65.48 +/- 24.7 % of predicted			
	69.03 +/- 24.0 % of predicted (type predose)	80.45 +/- 23.3 % of predicted	<0.0040	Absolute value, Tx vs. Ctl	

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.



Table 2-3. Symptom Score Outcomes

Citation	Study Arm	Number Evaluable	Treatment Duration (weeks)	Baseline Daytime Symptom Score	Final Daytime Symptom Score
Usual care vs. peak flow monitoring + action plan					
Jones, Mullee, Middleton et al. 1995	Usual care	64	45	26	4.95 (median; scale, 0-3)
	PFM-based action plan	63	39	26	2.85 (median; scale, 0-3)
Drummond, Abdalla, Beattie et al. 1994 (GRASSIC)	Usual care	284	67	52	
	PFM-based action plan	285	54	52	
Ayres, Campbell, Follows 1995	Usual care	64	64	24	1.91 +/- 0.6 (scale, 0-3)
	PFM-based action plan	61	61	24	1.77 +/- 0.6 (scale, 0-3)
Cowie, Revitt, Underwood et al. 1997	Usual care	48	48	24	
	PFM-based action plan	46	46	24	
Cote, Cartier, Robichaud et al. 1997	Usual care	54			
	PFM-based action plan	50			
Usual care + PFM use alone vs. usual care + PFM-based action plan					
Ignacio-Garcia and Gonzalez-Santos 1995	Usual care + PFM use	44	35	28	
	Usual care + PFM-based action plan	50	35	28	
Charlton, Antoniou, Atkinson et al. 1994	Usual care + PFM use	43	37	52	0.22 (median; scale, 0-3)
	Usual care + PFM-based action plan	48	42	52	0.26 (median; scale, 0-3)

	P-Value	Final Nighttime Symptom Score (scale, 0-3)	P-Value	Comments
		0.75 (median; scale, 0-3)		Symptom score across study was divided by number of days w/diary data X 28 to give a monthly rate; sx score day = cough; sx score night = waking at night; median wheeze = 5.46; shortness of breath = 7.88; asthma restricting normal daily activities = 0.0
	NS <sup>1</sup>	0.35 (median; scale, 0-3)	NS <sup>1</sup>	Symptom score across study was divided by number of days w/diary data X 28 to give a monthly rate; sx score day = cough; sx score night = wakenings at night; median wheeze = 4.39; shortness of breath = 6.50; asthma restricting normal daily activities = 0.17.
				Night and day sx score outcome is only from a subgroup of patients reporting variation in outcome; 112/246 never reported sleep disturbances; 15/246 reported that their sleep was disturbed every night.
				Night and day outcome is only from a subgroup of patients reporting variation in outcome, controlled for peak flow, FEV <sub>1</sub> , duration of asthma; 114/239 never reported sleep disturbances; 14/239 reported that their sleep was disturbed every night.
		0.69 +/- 0.13, (scale 0-3)		Sx score day = overall severity of asthma. Changes in: sleep disturbance scores 1.89 → 0.69; cough at rest 1.08 → 0.69; wheeze at rest was 1.25 → 0.67; difficulty breathing 1.47 → 0.96; cough with activity = 1.75 → 1.30.
	NS <sup>1</sup>	0.67 +/- 0.14 (scale, 0-3)		Sx score day = overall severity of asthma. Changes in: sleep disturbance scores 1.79 → 0.67; cough at rest 1.00 → 0.87; wheeze at rest was 0.97 → 0.74; difficulty breathing 1.41 → 0.85; cough with activity = 1.48 → 1.28. All comparisons in sx scores between groups NS.
				No significant differences in other indexes of asthma control, including waking with asthma, beta <sub>2</sub> -agonist use, or self-rating of asthma severity differed among the groups at 3 months or at 6 months after entry.
				No significant differences in other indexes of asthma control, including waking with asthma, beta <sub>2</sub> -agonist use, or self-rating of asthma severity among the groups at 3 months or at 6 months after entry.
				Nighttime symptoms = total nighttime awakenings over total study. (Values not reported by AHRQ)
				Nighttime symptoms = total nighttime awakenings over total study.
		0.25 (median; scale, 0-3)		Sx score day = wheeze day; Sx score night = wheeze night; daily score for activity restriction was 0.13.
	NS <sup>1</sup>	0.15 (median; scale, 0-3)	NS <sup>1</sup>	Sx score day = wheeze day; Sx score night = wheeze night; daily score for activity restriction was 0.06, p < 0.05 compared to control.

Source:

Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.

## References

- Abramson MJ, Bailey MJ, Couper FJ, Driver JS, Drummer OH, Forbes AB, McNeil JJ, Haydn Walters E; Victorian Asthma Mortality Study Group. Are asthma medications and management related to deaths from asthma? *Am J Respir Crit Care Med* 2001;163(1):12-8.
- Ayres JG, Campbell LM, Follows, RMA. A controlled assessment of an asthma self-management plan involving a budesonide dose regimen. *OPTIONS Research Group. Eur Respir J* 1996;9(5):886-92.
- Blue Cross and Blue Shield Association Technology Evaluation Center. Management of Chronic Asthma: Evidence Report/Technology Assessment Number 44. AHRQ Publication No. 01-EO44. Rockville, MD: Agency for Healthcare Research and Quality. September 2001.
- Charlton I, Antonion AG, Atkinson J, Campbell MJ, Chapman E, Mackintosh T, Shapira D. Asthma at the interface: bridging the gap between general practice and a district general hospital. *Arch Dis Child* 1994;70(4):313-8.
- Charlton I, Charlton G, Broomfield J, Mullee MA. Evaluation of peak flow and symptoms only self management plans for control of asthma in general practice. *BMJ* 1990;301(6765):1355-9.
- Cote J, Cartier A, Robichaud P, Boutin H, Malo JL, Rouleau M, Fillion A, Lavalee M, Krusky M, Boulet LP. Influence on asthma morbidity of asthma education programs based on self-management plans following treatment optimization. *Am J Respir Crit Care Med* 1997;155(5):1509-14.
- Cowie RL, Revitt SC, Underwood MF, Field SK. The effect of a peak flow-based action plan in the prevention of exacerbations of asthma. *Chest* 1997;112(6):1534-8.
- Drummond N, Abdalla M, Beattie JAG, et al. Effectiveness of routine self monitoring of peak flow in patients with asthma. *Grampian Asthma Study of Integrated Care (GRASSIC). BMJ* 1994;308(6928):564-7.
- Gibson PG, Coughlan J, Wilson AJ, Abramson M, Bauman A, Hensley MJ, Walters EH. Self-management education and regular practitioner review for adults with asthma. *Cochrane Database Syst Rev* 2000(2):CD001117. Review.
- Ignacio-Garcia JM, Gonzales-Santos P. Asthma self-management education program by home monitoring of peak expiratory flow. *Am J Respir Crit Care Med* 1995;151(2 Pt 1):353-9.
- Jones KP, Mullee MA, Middleton M, Chapman E, Holgate ST. Peak flow based asthma self-management: a randomized controlled study in general practice. *British Thoracic Society Research Committee. Thorax* 1995;50(8):851-7.